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(54) Title: MODIFIED POLYUNSATURATED FATTY ACIDS

(57) Abstract

The present invention provides polyunsaturated fatty acid compounds having antimalarial and/or neutrophil stimulatory activity. The polyunsaturated fatty acids contain 18-25 carbons and 1-6 double bonds and are characterized in that they have one or two substitutions selected from the group consisting of β oxa, γ oxa, β thia and γ thia. It is also preferred that the polyunsaturated fatty acid compound includes a further substitution selected from the group consisting of hydroxy, hydroperoxy, peroxy, carboxymethyl substitutions or attached to an amino acid. The invention also provides a method of producing an unsaturated oxa substituted fatty acid comprising reacting an unsaturated fatty acid alcohol with a carbene that is inserted in the OH bond of the alcohol. The invention further provides a method of treating inflammation with a composition comprising at least one hydroxy, hydroperoxy or peroxy derivative of a polyunsaturated fatty acid having a C₁₈₋₂₄ carbon chain and 1-6 cis or trans double bonds.

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MODIFIED POLYUNSATURATED FATTY ACIDS

The present invention relates to new polyunsaturated fatty acids having antimarial activity and/or neutrophil stimulatory activity. The present invention further relates to a group of modified polyunsaturated fatty acids which have the ability to suppress cytokine production and cytokine action. Such fatty acids have enhanced stability when compared to naturally occurring polyunsaturated fatty acids. The present invention further relates to compositions including the polyunsaturated fatty acids as the active ingredient and methods of anti-malarial, anti-infective or anti-inflammatory treatment or prevention involving the administration of this composition.

Over half of the world's population is at risk from malaria, with about 500 million acute infections and approximately 1 million deaths recorded each year. (Tropical Diseases Progress in International Research, 1987-1988, Ninth Programme Report, UNDP/World Bank/WHO, Geneva, 43-49; Stevenson MM Preface In: Stevenson MM, Ed. Malaria: Host responses to Infection, CRC Press, Inc). The use of antimarial drugs is associated with major problems because of increased resistance and toxic side-effects. Most currently used antimarials are unsuitable for use in children (most at risk of potentially fatal cerebral malaria), pregnant women and the aged.

Inflammation may be caused by bacteria, viruses and /or other infective agents, opportunistic infections (which may be consequent on an immunodepressed state, for example resulting from cancer or therapy, particularly cytotoxic drug therapy or radiotherapy), autoimmunity or otherwise. Septic shock is an illustration of a disease involving inflammation. Many of the clinical features of Gram-negative septic shock may be reproduced in animals by the administration of LPS to animals can prompt severe metabolic and physiological changes which can lead to death. Associated with the injection of LPS is the extensive production of pro-inflammatory cytokines such as tumour necrosis factor alpha (TNF α). Cachexia, which is characteristic of chronic exposure to TNF or interleukin-6, is a common symptom of advanced malignancy and severe infection. It is characterised by abnormal protein and glucose metabolism and body wasting. Chronic administration of TNF IL-1 in mice, rats and/or humans

cause anorexia, weight loss and depletion of body lipid and protein within 7 to 10 days (Ceraiini *et al.* 1985. Immunol. Lett. 11. 173; Fong *et al.* 1989 J. Exp. Med. 170. 1627. Moldawer *et al.* Am. J. Physiol.. 254 G450-G456. 1988; Fong *et al.* Am. J Physiol. 256. R659-R665 (1989); McCarthy *et al.* Am. J. Clin. Nature. 42. 1179-1182. 1982). TNF levels have been measured in patients with cancer and chronic disease associated with cachexia.

TNF α and IL-1, with their common functional activities such as pyrogenicity, somnogenicity and being mediators of inflammation, have been implicated in the pathology of other diseases associated with chronic inflammation apart from toxic shock and cancer-related cachexia. TNF has been detected in synovial fluid in patients with both rheumatoid and reactive arthritis and in the serum of patients with rheumatoid arthritis (Saxne *et al.* 1988. Arthrit. Rheumat. 31. 1041). Raised levels of TNF have been detected in renal transplant patients during acute rejection episodes (Maury and Teppo 1987. J. Exp. Med. 166. 1132). In animals, TNF has been shown to be involved in the pathogenesis of graft-versus-host disease in skin and gut following allogenic marrow transplantation.

Administration of a rabbit anti-murine TNF antibody was shown to prevent the histological changes associated with graft-versus-host disease and to reduce mortality (Piquet *et al.* 1987. J. Exp. Med. 166. 1220). TNF has also been shown to contribute significantly to the pathology of malaria (Clark *et al.* 1987. Am. J. Pathol. 129. 192-199). Further, elevated serum levels of TNF have been reported in malaria patients (Scuderi *et al.* 1986. Lancet 2. 1364-1365).

Multiple sclerosis (MS) is a chronic demyelinating disease of the central nervous system and is the commonest chronic neurological disease of young adults. The incidence of MS and its pattern of distribution have been unchanged for decades. The disease remains essentially untreatable.

MS usually affects multiple areas of white matter in the central nervous system (CNS), most frequently, the preterminal white matter, brainstem, spinal cord and the optic nerves. The primary process destroys myelin

sheaths and eventually kills oligodendrocytes creating the characteristic plaque of MS.

The early development of the plaque is characterised by the development of
5 perivascular inflammation followed by the migration of lymphocytes, plasma cells and macrophages into the lesion. This is followed by astrocyte gliosis and the attempts of demyelination by oligodendrocytes. The plaque is surrounded by lymphocytes. Anti-T cell agents such as anti-CD4 treatment are effective. Such agents inhibit the proliferation of T-cells.

10

Although the aetiology of MS is still unknown, the focus of research efforts that have led to plausible hypotheses have been those of immune dysregulation including autoimmunity and genetic predisposition, both of which may play a role in the actual development of disease. Both TNF β (lymphotoxin) and TNF α are thought to play a role in the pathophysiology.

20 Multiple immunological abnormalities are reproducibly found in patients in the acute stage of the disease. The synthesis of immunoglobulins, although normal in the periphery, is increased in the central nervous system and the antibodies produced have a characteristic banding pattern. The antigenic specificity of these antibodies is not known and it is unclear whether they have a role to play in the progression of the disease.

25 Various stressors known to activate the immune system such as viral infection or surgery can also produce an exacerbation of MS. Other activators such as γ -interferon produce similar effects when administered. In addition, immunosuppressive anti-inflammatory therapy with corticosteroids for example, can produce modest remission or at least palliation for short periods of time.

30

Myelopathy, a disorder of the spinal cord, can have many different aetiologies, most of which are mediated by inflammation, including the following:-

Neurosyphilis:

35 b_{12} or folate deficiency:
sarcoidosis:

transverse myelitis:

arachidonitis:

cervical spondylitis:

motor neuron disease:

5 neurofibromatosis:

spinal cord compression from tumour, disc or arthritis:

lupus erythematosus of the spinal cord; and

viral encephalomyelitis

- 10 Chronic inflammation or, as more commonly known, chronic immune system activation occurs in response to persistent antigen whose origin may be exogenous or may result from an autoimmune state. Such chronic inflammation results in local tissue destruction and depending upon the type of inflammation can result in systemic effects due to the sustained
- 15 production of inflammatory mediators. Such inflammatory mediators include the cytokines which are soluble mediators produced by activated lymphocytes and macrophages and effect cellular communication and physiological response. Chronic immune activation can occur as a result of infectious disease, such as chronic fatigue syndrome or toxic shock
- 20 syndrome or through autoimmune mechanisms resulting in such conditions as rheumatoid arthritis, inflammatory bowel disease, Crohns Disease and other diseases such as graft versus host disease.

Rheumatoid arthritis (Marrow *et al.*, I "Autoimmune Rheumatic Disease".

- 25 Blackwell Scientific Publ. Oxford, UK, Chapter 4 pp148-207 (1987)) is a disease characterised by chronic inflammation and erosion of joints that may affect up to 3% of the population, including children. Symptoms of rheumatoid arthritis include morning stiffness, swelling and pain upon motion in at least one joint and joint swelling. Non-specific symptoms
- 30 including lethargy, anorexia and weakness as well as fever and lymphadenopathy (characteristic of immune activation) may antedate joint involvement. Extra-articular manifestations of rheumatoid arthritis include vasculitis, cataracts, uveitis, interstitial fibrosis, pericarditis and myocarditis, peripheral neuropathy, myeloid deposits, chronic anaemia and
- 35 subcutaneous and pulmonary nodules.

Genetic factors and infectious agents including bacteria, fungi, mycoplasmas and viruses have been associated with the development of rheumatoid arthritis. Mild rheumatoid arthritis may be treated with non-steroidal anti-inflammatory drugs while severe cases require systemic corticosteroids, anti-5 metabolites or cytotoxic agents. Experimentally, anti-CD4 monoclonal antibodies and anti-TNF α antibodies have been used to treat rheumatoid arthritis (Horneff *et al.* Cytokine 3 266-267 (1991); Horneff *et al.* Arth. Rheum. 34 129-140 (1991) and Shoenfeld *et al.* Clin. Exp. Rheum. 9. 663-673 (1991). Williams *et al.* 1992 PNAS 89. 9784).

10 Inflammatory bowel disease (IBD) and Crohns disease are chronic inflammatory conditions that fulfil some of the criteria of an autoimmune disease (Snook. Gut 31 961-963 (1991)). Inflammation and tissue damage involves the recruitment and activation of neutrophils, macrophages and 15 lymphocytes (MacDermott *et al.* Adv. Immunol. 42 285-328 (1988)) which generate cytokines and proinflammatory molecules such as prostaglandins and leukotrienes (MacDermott. Mt. Sinai J. Med. 57 273-278 (1990)). As a result of chronic activation of immunocompetent cells, IL-1, IL-6 (Starter. Immunol. Res. 10 273-278 (1990) : Fiocchi. Immunol. Res. 10 239-246 (1991)) 20 and TNF α (MacDermott. Mt. Sinai J. Med. 57 273-278 (1990)) are all elevated in IBD and Crohns Disease patients.

Drugs used to treat IBD and Crohns Disease include anti-inflammatory agents such as sulphasalazine (5-ASA) corticosteroids, cyclosporin A and 25 azathioprine (Hanauer. Scand. J. Gastroenterol. 25 (Supl. 175) 97-106 (1990); Peppercorn. Annal. Intern. Med. 112 50-60 (1990)). Experimentally, anti-CD4 and anti-TNF monoclonal antibodies have been used to successfully treat ulcerative colitis (Emmerich *et al.* Lancet 338 570-571 (1991)).

30 Whilst a host may react against a genetically incompatible graft producing a host-versus-graft response, an immunocompetent graft (such as bone marrow or intestinal tissue) may react against the host resulting in graft-versus-host disease. These reactions are mediated by allogenic responses directed against a foreign MHC molecule and are mimicked *in vitro* by the mixed 35 lymphocyte reaction (MLR). Graft/host interactions result in chronic inflammation surrounding the grafted tissue with an increase in markers of

immune activation such as are seen in ADS (Grant. Immunol. Today 12 171-172 (1991)). Treatment of the graft/host interactions currently include either azathioprine, cyclosporin A or methylprednisone and, more recently.

rapamycin (Spekowski *et al.* Transplantation 53 258-264 (1992); Huber *et al.*

5 Bibliotheca Cardiologica. 43 103-110 (1988)). Monoclonal antibodies specific for CD3 (Wissing *et al.* Clin Exp Immunol. 83 333-337 (1991)), CD4 (Reinke *et al.* Lancet 338 702-703 (1991)) and TNF α have been used experimentally to inhibit graft/host reactions.

10 As mentioned above PUFAs have a range of useful biological activities (see for example International Patent Application Nos. WO 93/00084 and WO 95/00607 and the references cited therein). Unfortunately, due to their limited stability *in vivo*, PUFAs have not achieved widespread use as therapeutic agents. The present inventors have developed substituted

15 PUFAs which while retaining biological activity have increased stability *in vivo* i.e. slower metabolic turnover. These new polyunsaturated fatty acid (PUFA) compounds have direct antimalarial activity. In addition to their direct antimalarial activity, the novel PUFA activate human neutrophils causing release of granule contents, and exhibit synergy with TNF in the

20 production of superoxide. Activation of human neutrophils by the PUFA results in enhanced ability of these cells to kill malaria parasite (*P. falciparum*) within red blood cells and also the bacteria *Staphylococcus aureus*.

25 Further, the present inventors have also found that certain polyunsaturated fatty acids and novel polyunsaturated fatty acids and their hydroxy and hydroperoxy derivatives suppress production of cytokines.

These new PUFAs include at least one β oxa, β thia, γ oxa or γ thia

30 substitution. While saturated β -oxa fatty acids can be obtained using the standard procedure for ether synthesis, by reaction of alkyl halides with the dianions of α -hydroxy acids or by treating α -halo acids with deprotonated alcohols, the unsaturated β -oxa fatty acids of the present invention are not accessible using normal methods. Attempts to obtain the unsaturated

35 compounds in this manner lead only to decomposition products, resulting from undesirable side reactions at the olefinic and allylic carbons.

In a recent variation of the standard procedure, saturated β -oxa fatty acids have been obtained through nucleophilic substitution reactions under less vigorous conditions, by treating diazoacetates, activated by complexation

5 with boron trifluoride etherate, with alcohols. However, boron trifluoride etherate is known to cause isomerization of alkenes and it is therefore unsuitable for use in the synthesis of unsaturated β -oxa fatty acids.

10 The present inventors have now found that unsaturated β -oxa fatty acids can be obtained, in good yields, by insertion of carbenes in the O-H bond of alcohols. There is no complication from other carbene insertion reactions and, of particular significance, the olefinic moieties are unaffected under the reaction conditions.

15 The carbene can be generated from the corresponding diazo acetate or diazo alkane, by treatment with a catalyst such as a rhodium salt. Reaction of the carbene with the complementary alcohol, which is either a derivative of α -hydroxy acetic acid or an unsaturated fatty alcohol affords the unsaturated β -oxa fatty acid. In a preferred embodiment the alcohols are those obtained
20 by reduction of naturally occurring unsaturated fatty acids or the corresponding esters, and reaction with an ester of diazo acetic acid affords the unsaturated β -oxa fatty acid.

25 The present inventors have also shown that both β -oxa and β -thia substituted fatty acids are unable to undergo β oxidation. In addition certain of the novel compounds display other properties which differ from those of natural PUFA including enhanced solubility, varied oxidation reduction potentials and different charge and polarity.

30 Accordingly, in a first aspect the present invention consists in a polyunsaturated fatty acid compound having antimalarial and/or neutrophil stimulatory activity, the polyunsaturated fatty acid containing 18-25 carbons and 1-6 double bonds and wherein the polyunsaturated fatty acid compound has one or two substitutions selected from the group consisting of β oxa, γ oxa, β thia and γ thia.

In a preferred embodiment of the present invention the polyunsaturated fatty acid compound includes a further substitution selected from the group consisting of hydroxy, hydroperoxy, peroxy and carboxymethyl substitutions. In another embodiment the substituted fatty acid is

5 covalently attached to an amino acid.

In a further preferred embodiment of the present invention the polyunsaturated fatty acid compound contains 20-25 carbon atoms and 3-6 double bonds and is preferably an n-3 to n-6 fatty acid.

10

In another preferred embodiment of the present invention the polyunsaturated fatty acid compound is 21 carbons with 3-4 double bonds containing a β oxa or β thia substitution. 22 carbon atoms with 3-4 double bonds containing a γ thia or β oxa substitution. 23 carbons with 3-4 double bonds containing a β thia substitution. 24 carbons with 3-4 double bonds containing a γ thia substitution. 25 carbons with 3-6 double bonds containing a β oxa substitution. 25 carbons with 3-6 double bonds containing a β thia substitution. or 23 carbons. 3-6 double bonds. β thia and α -carboxymethyl group.

15

20 In yet another preferred embodiment of the present invention the polyunsaturated fatty acid compound has a ω hydroxy substitution.

25

In another preferred form of the invention the polyunsaturated fatty acid compound is attached to an amino acid. preferably an aspartic acid or glycine.

30

In a second aspect the present invention consists in a method of producing an unsaturated oxa substituted fatty acid comprising reacting an unsaturated fatty acid alcohol with a carbene such that the carbene is inserted in the O-H bond of the alcohol.

35

In a preferred embodiment of this aspect of the present invention the unsaturated fatty acid alcohol contains 18-25 carbon atoms and 1-6 double bonds.

In a further preferred embodiment of this aspect of the present invention the carbene is synthesized via rhodium acetate catalysed of a diazo compound.

In another preferred embodiment of this aspect of the present invention an
5 unsaturated β oxa substituted fatty acid is produced.

In a third aspect the present invention consists in a method of treating
inflammation in a subject. the method comprising administering to the
subject a therapeutically effective amount of an anti-inflamunatory
10 composition comprising at least one hydroxy. hydroperoxy. or peroxy
derivative of a polyunsaturated fatty acid having a carbon chain of 18 to 24
carbon atoms and having 1-6 cis or trans double bonds.

In a preferred embodiment of this aspect of the present invention the poly
15 unsaturated fatty acid may contain oxygen or sulphur atoms within the
carbon chains as oxa or thia derivatives.

In a further preferred embodiment of this aspect of the present invention the
polyunsaturated fatty acid is selected from the group consisting of the
20 C20:4n-6 (5. 8. 11. 14-eicosatetraenoc acid). C20:5n-3 (5. 8. 11. 14. 17-
eicosapentaenoic). C22:6n-3 (4. 7. 10. 13. 16. 19-docosahexaenoic acid) and
arachidonic acid.

Neutrophil/macrophage stimulatory agents may have application in the
25 treatment of other infections including *Candida* sp. *Trypanosoma*.
Schistosomiasis. *Tuberculosis*. viruses eg *herpes*. *Sindbis virus*. *Legionella*,
Listeriosis. *Pneumocystis*. *Pseudomonas*. They would also be useful as
adjunct therapy in immunocompromised individuals including those
undergoing cancer chemotherapy and transplant recipients and burns
30 patients. In addition others. so called normal individuals may also be
treated eg the aged. children under 2. alcoholics who are known to have poor
phagocytic cell activity.

In order that the nature of the present invention may be more clearly understood preferred forms therof will now be described with reference to the following examples and Figures in which:-

- 5 Fig 1a shows Arachidonic acid 5,8,11,14-Eicosatetraenoic acid
- Fig 1b shows 5-Hydroperoxy-6E,8Z,11Z,14Z-Eicosatetraenoic acid
- Fig 1c shows 9-Hydroperoxy-5Z,7E,11Z,14Z-Eicosatetraenoic acid
- Fig 1d shows 8-Hydroperoxy-5Z,9E,11Z,14Z-Eicosatetraenoic acid
- Fig 1e shows 12-Hydroperoxy-5Z,8Z,10E,14Z Eicosatetraenoic acid
- 10 Fig 1f shows 11-Hydroperoxy-5Z,8Z,12E,14Z Eicosatetraenoic acid
- Fig 1g shows 15-Hydroperoxy-5Z,8Z,11Z,13E Eicosatetraenoic acid
- Fig 1h shows 5-hydroxy-6E,8Z,11Z,14Z-Eicosatetraenoic acid
- Fig 1i shows 9-Hydroxy-5Z,7E,11Z,14Z-Eicosatetraenoic acid
- Fig 1j shows 8-Hydroxy-5Z,9E,11Z,14Z-Eicosatetraenoic acid
- 15 Fig 1k shows 12-Hydroxy-5Z,8Z,10E,14Z Eicosatetraenoic acid
- Fig 1l shows 11-Hydroxy-5Z,8Z,12E,14Z Eicosatetraenoic acid
- Fig 1m shows 15-Hydroxy-5Z,8Z,11Z,13E Eicosatetraenoic acid
- Figs 1n to Fig 1z show a range of substituted PUFAs in which Y is hydroxy, hydroperoxy or peroxy.
- 20 Fig 2a shows β -oxa 23:4 (n-6)
- Fig 2b shows β -oxa 21:3 (n-6)
- Fig 2c shows β -oxa 21:3 (n-3)
- Fig 2d shows β -oxa 25:6 (n-3)
- Fig 2e shows β -oxa 21:4 (n-3)
- 25 Fig 2f shows β -thia 23:4 (n-6)
- Fig 2g shows β -thia 21:3 (n-6)
- Fig 2h shows β -thia 21:3 (n-3)
- Fig 2i shows β -thia 25:6 (n-3)
- Fig 2j shows α -carboxymethyl- β -thia 23:4 (n-6)
- 30 Fig 2k shows γ -thia 24:4 (n-6)
- Fig 2l shows γ -thia 22:3 (n-6)
- Fig 2m shows γ -thia 22:3 (n-3)
- Fig 2n shows 16-OH- β -oxa 22:3 (n-6)
- Fig 2o shows 16-OH β -oxa 22:3 (n-3).

Fig 3 shows the effect of 20:4, 20:4 methyl ester (ME), 20:4 hydroperoxy (HP) and 20:4 hydroxy (H) on cytokine production by human peripheral blood mononuclear cells (PBMC) stimulated by the mitogen PHA.

20:4 hydroperoxy and 20:4 hydroxy inhibit cytokine production. both

5 macrophage-derived cytokines (TNF α) and T cell derived cytokines (lymphokines, interferon, IL-2).

Fig 4 shows the effect of 22:6, 22:6HP, 22:6H, 20:5, 20:5HP and 20:5H on PHA-induced cytokine production. These compounds inhibit *in vitro* production of macrophage and T cell-derived cytokines.

10 Fig 5 shows 20:4, 20:4 hydroperoxy (HP) and 20:4 hydroxy (OH) inhibit TNF α -induced expression of E-selection by human umbilical vein endothelial cells.

Fig 6 shows the fatty acid stimulated release from neutrophil azurophilic granules.

15 Fig 7 shows the fatty acid stimulated release from neutrophil specific granules.

Fig 8 shows the enhancement of neutrophil mediated killing of *Staph aureus* by fatty acids.

20 Fig 9 (a) shows the comparative abilities of arachidonic acid and β -oxa 23:4 (n-3) to undergo β oxidation

Fig 9 (b) shows that the compound does not inhibit β -oxidation of arachidonic acid and hence is unlikely to interfere with normal cellular metabolism.

Table 1 shows the direct antimarial activity of novel fatty acids against

25 chloroquine sensitive *P. falciparum*.

Table 2 shows the ability of the novel fatty acids to stimulate neutrophils to kill *P. falciparum*.

Table 3 shows the direct effect of novel fatty acids on survival of chloroquine resistant *P. falciparum*.

30 Table 4 shows inhibition of mitogen stimulated proliferation of peripheral blood mononuclear proliferation by fatty acids.

Table 5 shows inhibition of PHA stimulated TNF α production by fatty acids.

Table 6 shows inhibition of *Staph aureus* stimulated interferon production by fatty acids.

METHODS**DETECTION OF TNF-INDUCED ELAM-1 AND ICAM-1**

Human umbilical cord endothelial cells (5×10^4 , passage 1) were plated onto

- 5 the centre 60 wells of 96 well microtrays. the remaining wells were filled with pyrogen-free distilled water to reduce evaporation in the wells containing cells. The cells were fed with fresh medium (M199 supplemented with 20% FCS and endothelial cell growth supplement) every second day until confluent. On the day preceding the assay the cells were
- 10 washed with HBSS and re-fed. At the time of assay the medium was aspirated and fatty acids prepared in media were added to each well and incubated for various times as shown in Figs. At 37°C in 5% CO₂. After this incubation the cells were washed and treated with either TNF or PMA before being washed three times in PBS and fixed (100µl. 0.025%
- 15 glutaraldehyde in PBS. overnight at 4°C). The wells were washed with PBS and either anti-ELAM (Becton-Dickinson. 100µl. 1/1,000 in M199 containing 20% FCS) or anti-ICAM (Immunotech. 100µl. 12/2500 in M199 containing 20% FCS) was added and incubated for 1 hour at room temperature after which the wells were washed with PBS and 100µl of rabbit anti-mouse Ig-
- 20 HRP conjugate was added (1/1,000. Dako). A further incubation at room temperature for one hour followed before the plates were washed three times prior to the addition of 100µl of the substrate (ABTS). Colour development was measured at 410nm in a microplate reader.

25 MONONUCLEAR CELL PROLIFERATION ASSAYS

Mononuclear cells were separated from peripheral blood of normal human

donors as described by Ferrante and Thong (1978.....). The mononuclear

30 cells were resuspended in RPMI-1640 containing 20% human AB serum and placed into 96 well microtrays (50µl per well. cell density 4×10^6 cells/ml).

Fatty acid (66µm) was then added in 50µl and pre-incubated with the cells for 30 min at 37°C in 5% CO₂. Mitogen (PHA. ConA. PWM. Staph. aureus) was then added in 100µl and the cells incubated for 66 hours at 37°C in 5% CO₂ before the addition of tritiated thymidine (1µCi/well). After a total of

- 35 72h in culture the cells were harvested and proliferation (thymidine incorporation) and supernatant's assayed for the presence of cytokines.

CYTOKINE ASSAYS

5 Cytokine levels in culture supernatants were determined by specific ELISA using anti-cytokine antibodies. The following cytokine levels were determined TNF α . TNF β . interferon- γ . IL-1 β . IL-2.

PREPARATION OF FATTY ACID MICELLES AND PRETREATMENT OF NEUTROPHILS

10 To overcome fatty acid insolubility in aqueous solution mixed dipalmitoyl phosphatidylcholine (DPC. 400 μ g):fatty acid (100 μ g) micells were prepared in HBSS by sonication. Neutrophils were pretreated for 30 min at 37 $^{\circ}$ C. In some experiments PUFA were solubilized in ethanol.

15 MALARIA GROWTH INHIBITION ASSAY

20 Aliquots of RBC (5 \times 10 6 with 30% parasitaemia) was mixed with neutrophils (10 6) in the wells of a 96-well microtitre plate and then incubated for 2 hr at 37 $^{\circ}$ C in 5% CO $_2$ in air before adding 1 μ Ci of 3H-hypoxanthine. The plates were then incubated overnight. Individual well contents were collected onto glass filter paper and incorporated 3H measured in a liquid scintillation counter. Percent growth inhibition of the parasite was then calculated.

25 MEASUREMENT OF NEUTROPHIL CHEMILUMINESCENCE

30 To 100 μ l of neutrophils (1 \times 10 6) in HBSS was added 100 μ l of fatty acid micelles or DPC alone and an additional 300 μ l of HBSS. This was followed immediately by the addition of 500 μ l of lucigenin (0.25mg/ml in PBS and the resulting light output (mV) measured over time in a luminometer. Experiments were performed in triplicate with cells from a separate individual and values presented represent peak values of the responses.

MEASUREMENT OF DEGRANULATION

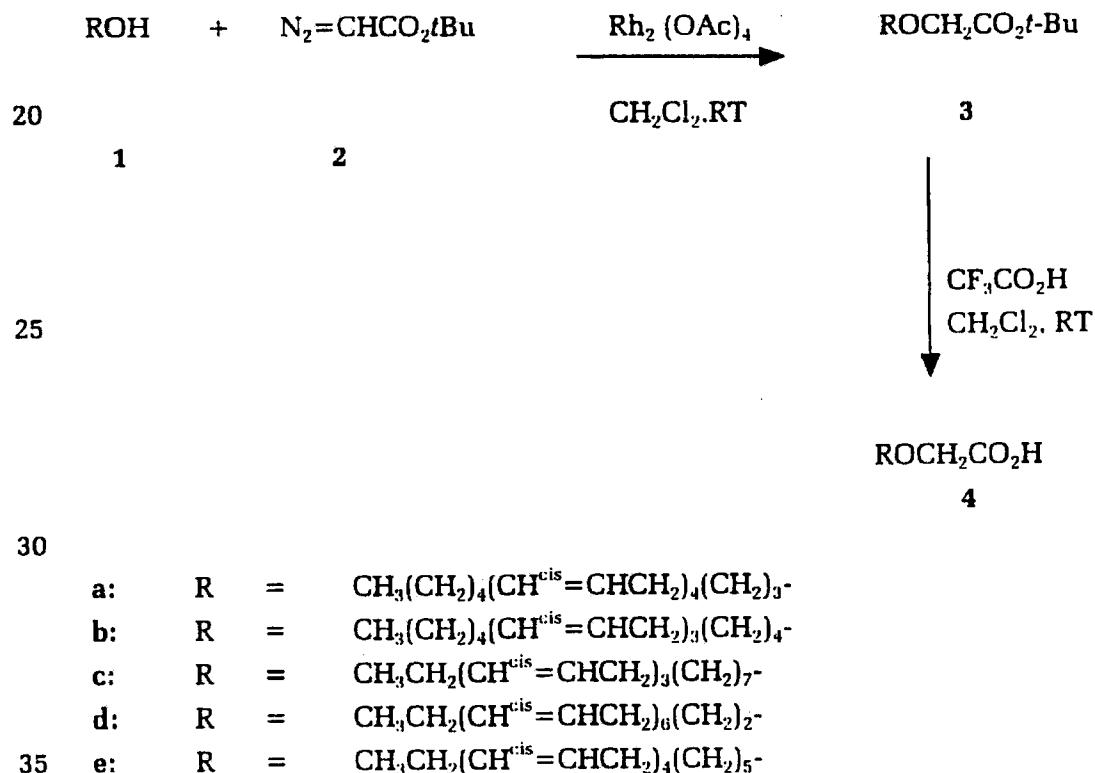
Degranulation was determined by measuring vitamin B12 binding protein (as described by Gottlieb *et al.* 1965. Blood 25:875-883) and β -glucuronidase release (as described by Kolodenev and Mumford. 1976. Clin. Chem. Acta 70:247-257).

BACTERICIDAL ASSAY

10 Neutrophil bactericidal activity against *Staphylococcus aureus* was measured according to the procedure described by Ferrante and Abell. 1986. Infect. Immun. 51:607.

15 CHEMICAL SYNTHESIS

Synthesis of β -Oxa Fatty Acids



REAGENTS

Arachidonyl Alcohol (**1a**) - Nu Chek Prep.. Elysian. MN. USA
Gammma Linolenyl Alcohol (**1b**) - Nu Chek Prep.. Elysian. MN. USA
5 Linolenyl Alcohol (**1c**) - Nu Chek Prep.. Elysina. MN. USA
Docosahexaenyl Alcohol (**1d**) - Nu Chek Prep.. Elysian. MN. USA
Z.Z.Z.Z- 6. 9. 12. 15-Octadecatetraenyl alcohol (**1e**) - Synthesised by lithium
aluminium hydride reduction of Methyl 6. 9. 12. 15-Octadecatetraenoate.
Methyl Z.Z.Z.Z- 6. 9. 12. 15-Octadecatetraenoate - Sigma Chemical Company
10 Rhodium Acetate Dimer - Aldrich Chemical Company
tert-Butyl DiazoAcetate (**2**) - synthesized from *tert*-Butyl Acetoacetate as per:
Regitz. M: Hocker. J: Leidhegener. A. *Organic Syntheses Coll. Vol. 5.* 179.
tert-Butyl Acetoacetate - Fluka AG
Trifluoroacetic Acid - Aldrich Chemical Company.
15 All solvents were distilled prior to use.
Column chromatographies were performed under positive nitrogen pressure
on Merck Silica Gel 60 (230-400 mesh). Art. 9385.

20 PROCEDURE

tert-Butyl alkyloxyacetates **3**
The relevant fatty alcohol **1** (1 mol equivalent) was weighed into a two-neck
round bottom flask under dry nitrogen and was dissolved in
dichloromethane. To this stirred solution was added rhodium acetate dimer
25 (0.5 mol%). followed by the dropwise addition of a solution of *tert*-butyl
diazoacetate **2**(2.5 mol equivalents) in dichloromethane via syringe. After
the addition was complete the reaction mixture was stirred at room
temperature under nitrogen for 2 hrs. The crude reaction mixture was
concentrated under a stream of dry nitrogen and the residue was purified by
30 column chromatography on silica, eluting with hexane/diethyl ether (9:1), to
afford the relevant *tert*-butyl alkloxyacetate **3** as an oil.

ALKYLOXYACETIC ACIDS **4**

35 The relevant *tert*-butyl alkyloxyacetate **3** (ca. 100mg. 1 mol equivalent) was
weighed into a two-neck round-bottom flask under dry nitrogen and was

dissolved by the addition of dichloromethane (ca. 4ml). To this stirred solution was added trifluoroacetic acid (ca. 1ml), and the reaction mixture was stirred at room temperature under nitrogen for 2 hrs. The crude reaction mixture was concentrated under a stream of dry nitrogen and the residue

5 was purified by column chromatography on silica, eluting with hexane/diethyl ether/acetic acid (40:60:2), to afford the relevant alkyloxyacetic acid **4** as an oil.

t-Butyl (5, 8, 11, 14-eicosatetraenyoxy) acetate (3a)

10 [t-Bu β -oxa 23:4 (n-6)]

^1H n.m.r. (200MHz. CDCl_3) δ 0.89 (3H. t. J 6.7 Hz. $\text{C}20'$ - H_3 , 1.25 - 1.38 (8H. m. $\text{C}3'$ - H_2 , $\text{C}17'$ - H_2 , $\text{C}18'$ - H_2 , $\text{C}19'$ - H_2), 1.49 (9H. s. $\text{C}(\text{CH}_3)_3$), 1.56 - 1.69 (2H. m. $\text{C}2'$ - H_2), 2.01 - 2.15 (4H. m. $\text{C}4'$ - H_2 , $\text{C}16'$ - H_2), 2.79 - 2.87 (6H. m. $\text{C}7'$ - H_2 , $\text{C}10'$ - H_2 , $\text{C}13'$ - H_2), 3.52 (2H. t. J 6.6 Hz. $\text{C}1'$ - H_2), 3.94 (2H. s. $\text{C}2$ - H_2), 5.32 -

15 5.45 (8H. m. $\text{C}5'$ - H , $\text{C}6'$ - H , $\text{C}8'$ - H , $\text{C}9'$ - H , $\text{C}11'$ - H , $\text{C}12'$ - H , $\text{C}14'$ - H , $\text{C}15'$ - H);

^{13}C n.m.r. (50MHz. CDCl_3) δ 169.82s. 130.48d. 129.97d. 128.55d.

128.42d. 128.08d. 128.02d. 127.96d. 127.59d. 81.40s. 71.63t. 68.83t. 31.58t, 29.29t. 29.07t. 28.14q. 27.23t. 27.01t. 26.09t. 25.66t. 22.57t. 14.09q.

20 **t-Butyl Z,Z,Z-(6, 9, 12-octadecatrienyoxy) acetate (3b)**

[t-Bu β -oxa 21:3(n-6)]

^1H n.m.r. (200 MHz. CDCl_3) δ 0.89 (3H. t. J 6.7 Hz. $\text{C}18'$ - H_3 , 1.25 -

1.45 (10H. m. $\text{C}3'$ - H_2 , $\text{C}4'$ - H_2 , $\text{C}15'$ - H_2 , $\text{C}16'$ - H_2 , $\text{C}17'$ - H_2), 1.48 (9H. s. $\text{C}(\text{CH}_3)_3$), 1.56 - 1.68 (2H. m. $\text{C}2'$ - H_2), 2.01 - 2.13 (4H. m. $\text{C}5'$ - H_2 , $\text{C}14'$ - H_2),

25 2.77 - 2.84 (4H. m. $\text{C}8'$ - H_2 , $\text{C}11'$ - H_2), 3.51 (2H. t. J 6.6 Hz. $\text{C}1'$ - H_2), 3.94 (2H. s. $\text{C}2$ - H_2), 5.27 - 5.48 (6H. m. $\text{C}6'$ - H , $\text{C}7'$ - H , $\text{C}9'$ - H , $\text{C}10'$ - H , $\text{C}12'$ - H , $\text{C}13'$ - H);

^{13}C n.m.r. (50MHz. CDCl_3) δ 169.82s. 130.41d. 130.10d. 128.33d.

128.21d. 127.87d. 127.64d. 81.39s. 71.72t. 68.82t. 31.52t. 29.57t. 29.49t, 28.12q. 27.18t. 25.74t. 25.63t. 22.56t. 14.04q.

30

t-Butyl Z,Z,Z-(9, 12, 15-octadecatrienyoxy)acetate (3c)

[t-Bu β -oxa 21:3(n-3)]

^1H n.m.r. (200 MHz. CDCl_3) δ 0.98 (3H. t. J 7.5 Hz. $\text{C}18'$ - H_3 , 1.25 -

1.40 (10H. m. $\text{C}3'$ - H_2 , $\text{C}4'$ - H_2 , $\text{C}5'$ - H_2 , $\text{C}6'$ - H_2 , $\text{C}7'$ - H_2), 1.48 (9H. s. $\text{C}(\text{CH}_3)_3$),

35 1.51 - 1.67 (2H. m. $\text{C}2'$ - H_2), 2.01 - 2.15 (4H. m. $\text{C}8'$ - H_2 , $\text{C}17'$ - H_2), 2.75 - 2.86

(4H. m. C11'-H₂, C14'-H₂), 3.50 (2H. t. J6.6 Hz. C1'-H₂), 3.95 (2H. s. C2-H₂), 5.31 - 5.43 (6H. m. C9'-H. C10'-H. C12'-H. C13'-H. C15'-H. C16'-H);
¹³C n.m.r. (50MHz. CDCl₃) δ 169.84s. 131.94d. 130.35d. 128.27d. 127.66d. 127.13d. 81.36s. 71.83t. 68.82t. 29.64t. 29.45t. 29.25t. 28.12q.
5 27.24t. 26.04t. 25.62t. 25.53t. 20.53t. 14.25q.

t-Butyl Z,Z,Z,Z,Z-(4,7,10,13,16,19-docosahexaenoxy)acetate (3d)

[t-Bu B-oxa 25:6(n-3)]

10 ¹H n.m.r. (200 MHz. CDCl₃) δ 0.98 (3H. t. J 7.5 Hz. C22'-H₃), 1.48 (9H. s. C(CH₃)₃), 1.58 - 1.76 (2H. m. C2'-H₂), 2.00 - 2.21 (4H. m. C3'-H₂, C21'-H₂), 2.79 - 2.87 (10H. m. C6'-H₂, C9'-H₂, C12'-H₂, C15'-H₂, C18'-H₂), 3.52 (2H. t. J 6.6 Hz. C1'-H₂), 3.95 (2H. s. C2-H₂), 5.28-5.46 (12H. m. C4'-H. C5'-H. C7'-H. C8'-H. C10'-H. C11'-H. C13'-H. C14'-H. C16'-H. C17'-H. C19'-H. C20'-H);
¹³C n.m.r. (50MHz. CDCl₃) δ 169.76s. 132.03d. 129.41d. 128.57d. 128.39d. 128.36d. 128.24d. 128.21d. 128.16d. 128.12d. 128.02d. 127.88d. 127.02d. 81.40s. 71.08t. 68.83t. 29.53t. 28.12q. 25.63t. 25.59t. 25.54t. 23.72t. 20.55t. 14.25q.

t-Butyl Z,Z,Z,Z-(6, 9, 12, 15-octadecatetraenoxy acetate (3e)

20 [t/Bu β-oxa 21:4(n-3)]

10 ¹H n.m.r. (200 MHz. CDCl₃) δ 0.98 (3H. t. J 7.5 Hz. C18'-H₃), 1.21 - 1.52 (4H. m. C3'-H₂), 1.48 (9H. s. C(CH₃)₃), 1.53 - 1.66 (2H. m. C2'-H₂), 2.01 - 2.15 (4H. m. C5'-H₂, C17'-H₂), 2.77 - 2.87 (6H. m. C8'-H₂, C11'-H₂, C14'-H₂), 3.51 (3H. t. J6.6 Hz. C1'-H₂), 3.95 (2H. s. C2-H₂), 5.28 - 5.47 (8H. m. C6'-H. C7'-H. C9'-H. C10'-H. C12'-H. C13'-H. C15'-H. C16'-H);
¹³C n.m.r. (50MHz. CDCl₃) δ 169.82s. 132.00d. 130.15d. 128.49d. 128.42d. 128.01d. 127.96d. 127.78d. 127.05d. 81.39s. 71.72t. 68.81t. 29.57t. 29.49t. 28.12q. 27.18t. 25.74t. 25.63t. 25.54t. 20.55t. 14.25q.

30 **Z,Z,Z,Z-(5, 8, 11, 14-Eicosatetraenoxy) acetic acid (4a)**

[β-oxa 23:4(n-3)]

10 ¹H n.m.r. (200 MHz. CDCl₃) δ 0.89 (3H. t. J 6.6 Hz. C20'-H₃), 1.25 - 1.490 (8H. m. C3'-H₂, C17'-H₂, C18'-H₂, C19'-H₂), 1.57 - 1.74 (2H. m. C2'-H₂), 2.00 - 2.14 (4H. m. C4'-H₂, C16'-H₂), 2.78 - 2.85 (6H. m. C7'-H₂, C10'-H₂, C13'-H₂), 3.58 (2H. br t. J6.0 Hz. C1'-H₂), 4.08 (2H. s. C2-H₂), 5.29 - 5.46 (8H. m. C5'-H. C6'-H. C8'-H. C9'-H. C11'-H. C12'-H. C14'-H. C15'-H);
35

¹³C n.m.r. (50MHz. CDCl₃) δ 173.37s. 130.47d. 129.64d. 128.55d. 128.24d. 128.13d. 128.02d. 127.86d. 127.152d. 71.81t. 31.49t. 29.49t. 29.21t. 27.20t. 26.88t. 26.03t. 25.63t. 25.63t. 22.54t. 14.02q.

5 Z,Z,Z-(6, 9, 12-Octadecatrienyloxy) acetic acid (4b)

[β-oxa 21:3(n-6)]

¹H n.m.r. (200 MHz. CDCl₃) δ 0.89 (3H. t. J 6.8 Hz. C18'-H₃), 1.23 - 1.43 (10H. m. C3'-H₂, C4'-H₂, C15'-H₂, C16'-H₂, C17'-H₂), 1.51 - 1.71 (2H. m. C2'-H₂), 2.00 - 2.10 (4H. m. C5' - H₂, C14'-H₂), 2.75 - 2.86 (4H. m. C8'-H₂, C11'-H₂), 3.60 (2H. t. J 6.6 Hz. C1'-H₂) 4.17 (2H. s. C2-H₂), 5.26 - 5.47 (6H. m. C6'-H, C7'-H, C9'-H, C10'-H, C12'-H, C13'-H)

¹³C n.m.r. (50MHz. CDCl₃) δ 171.74s. 130.45d. 129.84d. 128.40d. 128.20d. 128.10d. 127.59d. 71.13d. 31.51d. 29.36d. 29.31d. 27.20d. 27.10d. 25.62d. 22.57d. 22.55d. 14.04q.

15

Z,Z,Z-(9, 12, 15-Octadecatrienyloxy) acetic acid (4c)

[β-oxa 21:3(n-3)]

¹H n.m.r. (200 MHz. CDCl₃) δ 0.97 (3H. t. J 7.5 Hz. C18'-H₃), 1.25 - 1.43 (10H. m. C3'-H₂, C4'-H₂, C5'-H₂, C6'-H₂, C7'-H₂), 1.51 - 1.63 (2H. m. C2'-H₂), 2.01 - 2.15 (4H. m. C8'-H₂, C17'-H₂), 2.75 - 2.86 (4H. m. C11' - H₂, C14'-H₂), 3.55 (2H. t. J 6.5 Hz. C1'-H₂), 4.12 (2H. s. C2'-H₂), 5.28 - 5.46 (6H. m. C9'-H, C10'-H, C12'-H, C13'-H, C15'-H, C16'-H)

¹³C n.m.r. (50MHz. CDCl₃) δ 174.73s. 131.93d. 130.29d. 128.25d. 127.68d. 127.11d. 72.16t. 29.60t. 29.41t. 29.20t. 27.21t. 25.88t. 25.60t. 25.52t.

25 20.53t. 14.24q.

Z,Z,Z,Z,Z,Z-(4, 7, 10, 13, 16, 19-Docosahexaenyloxy) acetic acid (4d)

[β-oxa 25:6(n-3)]

¹H n.m.r. (200 MHz. CDCl₃) δ 0.97 (3H. t. J 7.5 Hz. C22'-H₃), 1.65 - 1.78 (2H. m. C2'-H₂), 2.01 - 2.21 (4H. m. C3'-H₂, C21'-H₂), 2.75 - 2.92 (10H. m. C6' - H₂, C9'-H₂, C12'-H₂, C15'-H₂, C18'-H₂), 3.57 (2H. t. J 6.4 Hz. C1'-H₂), 4.12 (2H. s. C2-H₂), 5.28 - 5.46 (12H. m. C4'-H, C5'-H, C7'-H, C8'-H, C10'-H, C11'-H, C13'-H, C14'-H, C16'-H, C17'-H, C19'-H, C20'-H), 10.22 (1H. br. CO₂H)

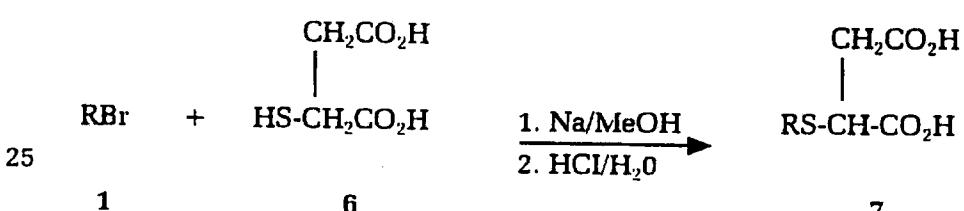
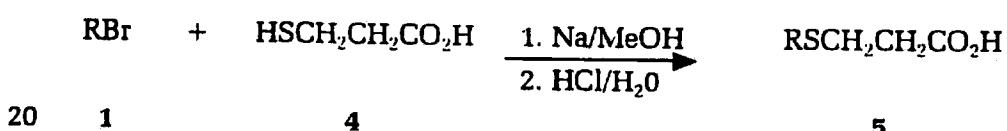
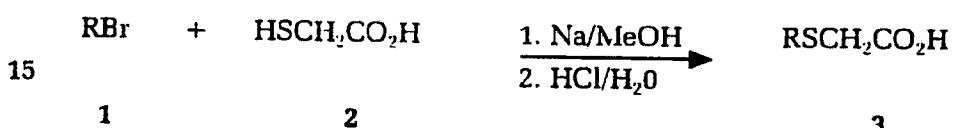
¹³C n.m.r. (50MHz. CDCl₃) δ 172.99s. 132.04d. 128.98d. 128.75d. 128.59d. 128.49d. 128.38d. 128.28d. 128.22d. 128.17d. 128.08d. 127.86d. 127.01d. 71.48t. 29.26t. 25.64t. 25.59t. 25.54t. 23.55t. 14.25q.

Z,Z,Z,Z-(6, 9, 12, 15-Octadecatetraenyl) acetic acid (4e)

[β -oxa 21:4(n-3)]

¹H n.m.r. (200 MHz, CDCl_3) δ 0.97 (3H, t, J 7.5 Hz, $\text{C}18'$ -H₃), 1.33 - 5 1.40 (4H, m, $\text{C}3'$ -H₂, $\text{C}4'$ -H₂), 1.54 - 1.68 (2H, m, $\text{C}2'$ -H₂), 2.00 - 2.15 (4H, m, $\text{C}5'$ - H₂, $\text{C}17'$ -H₂), 2.77 - 2.87 (6H, m, $\text{C}8'$ -H₂, $\text{C}11'$ -H₂, $\text{C}14'$ -H₂), 3.56 (3H, t, J 6.6 Hz, $\text{C}1'$ -H₂), 4.11 (2H, s, $\text{C}2$ -H₂), 5.24 - 5.45 (8H, m, $\text{C}6'$ -H, $\text{C}7'$ -H, $\text{C}9'$ -H, $\text{C}10'$ -H, $\text{C}12'$ -H, $\text{C}13'$ -H, $\text{C}15'$ -H, $\text{C}16'$ -H);
¹³C n.m.r. (50MHz, CDCl_3) δ 172.05s, 132.03d, 139.92d, 128.53d, 10 128.35d, 128.08d, 128.01d, 127.94d, 127.04d, 71.10t, 29.34t, 27.10t, 25.64t, 25.55t, 20.55t, 14.25g.

Synthesis of β and γ Thia fatty acids.



a: $R = \text{CH}_3(\text{CH}_2)_4(\text{CH}^{cis}=\text{CHCH}_2)_2(\text{CH}_2)_3$

b: R = $\text{CH}_2(\text{CH}_2)_4(\text{CH}^{\text{cis}}=\text{CHCHCH}_2)_2(\text{CH}_2)_4$

30 c: R = $\text{CH}_2\text{CH}_2(\text{CH}_2^{\text{cis}}=\text{CHCH}_2)_n(\text{CH}_2)_m$

d: $\text{R} = \text{CH}_2\text{CH}_2(\text{CH}^{\text{cis}}=\text{CHCH}_2)_2(\text{CH}_2)_2$

Reagents

Fatty bromides (1a-1d) - synthesized from the corresponding fatty alcohols by treatment with triphenylphosphine and carbon tetrabromide in dichloromethane.

5 *Arachidonyl alcohol* - Nu Chek Prep.

Gamma linolenyl alcohol - Nu Chek Prep

Linolenyl alcohol - Nu Check Preo

Docosahexaenyl alcohol - Nu Chek Prep

Mercaptoacetic acid - Aldrich Chemical Company

10 *Mercaptopropionic acid* - Aldrich Chemical Company

All solvents were distilled prior to use.

Column chromatographies were performed under positive nitrogen pressure on Merck Silica Gel 60 (230-400mesh).Art. 9385.

15 **PROCEDURE**

Alkylthioacetic acids 3a-d

Sodium (3mol equivalents) was dissolved in methanol in a two-neck round-bottomed flask under dry nitrogen and to this stirred solution was added mercaptoacetic acid (1.2 mol equivalents). After the initial white precipitate

20 had dissolved, a solution of the relevant bromide 1 (1 mol equivalent) in diethyl ether was added via syringe and the mixture was stirred at room temperature under nitrogen for 16hr. The crude reaction mixture was poured into an equal volume of hydrochloric acid (10% v/v) and extracted with diethyl ether. The resulting extract was concentrated under a stream of dry nitrogen and the residue was purified by flash chromatography on silica, eluting with hexane/diethyl ether/acetic acid (40:60:2) to afford the relevant alkylthioacetic acid 3 as an oil.

Alkylthiopropionic acids 5a-c

30 The alkylthiopropionic acids 5a-c were synthesized by alkaline condensation of the respective fatty bromides 1a-c with mercaptopropionic acid 4, in an analogous manner to that described above for the alkylthioacetic acids 3a-d.

Z,Z,Z,Z-(5,8,11,14-Eicosatetraenylthio)succinic Acid 7a

(5,8,11,14-Eicosatetraenylthio)succinic acid 7a was synthesized by condensation of the fatty bromide 1a (1 mol equivalent) with mercaptosuccinic acid 6 (1.2 mol equivalents), in the presence of sodium

5 (4.5 mol equivalents) in an analogous manner to that described above for the alkylthioacetic acids 3a-d.

Z,Z,Z,Z-(5,8,11,14-Eicosatetraenylthio) acetic acid (3a)

[β -thia 23:4(n-6)]

10 ^1H n.m.r. (200 MHz, CDCl_3) δ 0.89 (3H, t, J 6.7 Hz, $\text{C}20'$ - H_3), 1.21 - 1.54 (8H, m, $\text{C}3'$ - H_2 , $\text{C}17'$ - H_2 , $\text{C}18'$ - H_2 , $\text{C}19'$ - H_2), 1.57 - 1.72 (2H, m, $\text{C}2'$ - H_2), 2.01 - 2.14 (4H, m, $\text{C}4'$ - H_2 , $\text{C}16'$ - H_2), 2.67 (2H, t, J 7.2 Hz, $\text{C}1'$ - H_2), 2.73 - 2.88 (6H, m, $\text{C}7'$ - H_2 , $\text{C}10'$ - H_2 , $\text{C}13'$ - H_2), 3.25 (2H, s, $\text{C}2$ - H_2), 5.27 - 5.47 (8H, m, $\text{C}5'$ - H , $\text{C}6'$ - H , $\text{C}8'$ - H , $\text{C}9'$ - H , $\text{C}11'$ - H , $\text{C}12'$ - H , $\text{C}14'$ - H , $\text{C}15'$ - H);
 15 ^{13}C n.m.r. (50MHz, CDCl_3) δ 175.69s, 130.49d, 129.54d, 128.56d, 128.27d, 128.15d, 127.89d, 127.55d, 33.45t, 32.70t, 31.51t, 30.38t, 29.32t, 28.62t, 28.46t, 27.23t, 26.73t, 25.65t, 22.57t, 14.05q.

Z,Z,Z,Z-(6,9, 12-Octadecatrienylthio)acetic acid (3b)

20 [β -thia 21:3(n-6)]

^1H n.m.r. (200 MHz, CDCl_3) δ 0.89 (3H, t, J 6.6 Hz, $\text{C}18'$ - H_3), 1.23 - 1.48 (10H, m, $\text{C}3'$ - H_2 , $\text{C}4'$ - H_2 , $\text{C}15'$ - H_2 , $\text{C}16'$ - H_2 , $\text{C}17'$ - H_2), 1.52 - 1.70 (2H, m, $\text{C}2'$ - H_2), 1.89 - 2.15 (4H, m, $\text{C}5'$ - H_2 , $\text{C}14'$ - H_2), 2.63 (2H, t, J 7.0 Hz, $\text{C}1'$ - H_2), 2.70 - 2.87 (4H, m, $\text{C}8'$ - H_2 , $\text{C}11'$ - H_2), 3.26 (2H, s, $\text{C}2$ - H_2), 5.29 - 5.47 (6H, m, $\text{C}6'$ - H , $\text{C}7'$ - H , $\text{C}9'$ - H , $\text{C}10'$ - H , $\text{C}12'$ - H , $\text{C}13'$ - H);
 ^{13}C n.m.r. (50MHz, CDCl_3) δ 176.54s, 130.41d, 129.86d, 128.367d, 128.11d, 128.03d, 128.59d, 33.49t, 32.74t, 31.51t, 30.36t, 29.31t, 29.14t, 28.81t, 28.34t, 27.21t, 27.05t, 25.63t, 22.55t, 14.04q.

Z,Z,Z-(9,12,15-Octadecatrienylthio)acetic acid (3c)

30 [β -thia 21:3(n-3)]

^1H n.m.r. (200 MHz, CDCl_3) δ 0.89 (3H, t, J 6.7 Hz, $\text{C}18'$ - H_3), 1.21 - 1.52 (10H, m, $\text{C}3'$ - H_2 , $\text{C}4'$ - H_2 , $\text{C}5'$ - H_2 , $\text{C}6'$ - H_2 , $\text{C}7'$ - H_2), 1.54 - 1.72 (2H, m, $\text{C}2'$ - H_2), 2.01 - 2.15 (4H, m, $\text{C}8'$ - H_2 , $\text{C}17'$ - H_2), 2.67 (2H, t, J 7.2 Hz, $\text{C}1'$ - H_2), 2.73 - 2.87 (4H, m, $\text{C}11'$ - H_2 , $\text{C}14'$ - H_2), 3.25 (2H, s, $\text{C}2$ - H_2), 5.27 - 5.48 (6H, m, $\text{C}9'$ - H , $\text{C}10'$ - H , $\text{C}12'$ - H , $\text{C}13'$ - H , $\text{C}15'$ - H , $\text{C}16'$ - H);

¹³C n.m.r. (50MHz, CDCl₃) δ 176.21s, 131.94d, 130.29d, 128.26d, 127.69d, 127.11d, 33.49t, 32.82t, 30.39t, 29.60t, 29.36t, 29.20t, 29.12t, 28.89t, 28.70t, 27.21t, 25.61t, 25.52t, 20.53t, 14.25q.

5 **Z,Z,Z,Z,Z-(9,12,15-Octadecatrienylthio)acetic acid (3d)**
 [β-thia 25:6(n-3)]
¹H n.m.r. (200 MHz, CDCl₃) δ 0.99 (3H, t, J 7.5 Hz, C22'-H₃), 1.62 - 1.77 (2H, m, C2'-H₂), 2.02 - 2.24 (4H, m, C3'-H₂, C21'-H₂), 2.68 (2H, t, J 7.4 Hz, C1'-H₂), 2.83 - 2.85 (10H, m, C6'-H₂, C9'-H₂, C12'-H₂, C15'-H₂, C18'-H₂),
 10 3.26 (2H, s, C2-H₂), 5.29 - 5.47 (12H, m, C4'-H₂, C5'-H, C7'-H, C8'-H, C10'-H, C11'-H, C13'-H, C14'-H, C16'-H, C17'-H, C19'-H, C20'-H);
¹³C n.m.r. (50MHz, CDCl₃) δ 176.37s, 132.01d, 128.94d, 128.68d, 128.55d, 128.24d, 128.18d, 128.12d, 128.07d, 127.85d, 126.99d, 33.41t, 32.21t, 28.67t, 26.12t, 25.64t, 25.22t, 20.53t, 14.24q.

15 **Z,Z,Z,Z-(5,8,11,14-Eicosatetraenylthio) propionic acid (5a)**
 [γ-thia 24:4(n-6)]
¹H n.m.r. (300 MHz, CDCl₃) δ 0.89 (3H, t, J 6.8 Hz, C20'-H₃), 1.26 - 1.38 (6H, m, C17'-H₂, C18'-H₂, C19'-H₂), 1.41 - 1.51 (2H, m, C3'-H₂), 1.56 - 1.66 (2H, m, C2' - H₂), 2.02 - 2.12 (4H, m, C4'-H₂, C16'-H₂), 2.54 (2H, t, J 7.3 Hz, C1'-H₂), 2.66 (2H, t, J 6.6 Hz C3-H₂), 2.78 (2H, t, J 6.6 Hz, C2-H₂), 2.78 - 2.86 (6H, m, C7'-H₂, C10'-H₂, C13'-H₂), 5.29 - 5.44 (8H, m, C5'-H, C6'-H, C8'-H, C9'-H, C11'-H, C12'-H, C14'-H, C15'-H);
¹³C n.m.r. (50MHz, CDCl₃) δ 178.22s, 130.37d, 129.58d, 128.46d, 128.19d, 128.10d, 128.03d, 127.81d, 127.48d, 34.68t, 32.046t, 31.44t, 29.25t, 29.01t, 28.67t, 27.15t, 26.70t, 26.54t, 25.57t, 22.50t, 13.98q.

30 **Z,Z,Z-(6,9,12-Octadecatrienylthio)propionic acid (5b)**
 [γ-thia 22:3(n-6)]
¹H n.m.r. (200 MHz, CDCl₃) δ 0.89 (3H, t, J 6.8 Hz, C18'-H₃), 1.25 - 1.42 (10H, m, C3'-H₂, C4'-H₂, C15'-H₂, C16'-H₂, C17'-H₂), 1.53 - 1.65 (2H, m, C2'-H₂), 2.01 - 2.10 (4H, m, C5' - H₂, C14'-H₂), 2.53 (2H, t, J 7.3 Hz, C1'-H₂), 2.66 (2H, t, J 6.8 Hz, C3-H₂), 2.78 (2H, t, J 6.8 Hz C2-H₂), 2.74 - 2.83 (4H, m, C8'-H, C11'-H), 5.26 - 5.47 (6H, m, C6'-H, C7'-H, C9'-H, C10'-H, C12'-H, C13'-H);

35

¹³C n.m.r. (50MHz. CDCl₃) δ 178.14s. 130.38d. 129.90d. 128.33d. 128.10d. 127.97d. 127.58D. 34.69t. 32.13t. 31.49t. 29.41t. 29.29t. 29.18t. 28.46t. 27.19t. 27.06t. 26.58t. 25.61t. 14.03q.

5 **Z,Z,Z-(9,12,15-Octadecatrienylthio)proponionic acid (5c)**
 [γ-thia 22:3(n-3)]
¹H n.m.r. (200 MHz. CDCl₃) δ 0.98 (3H. t, J 7.5 Hz. C18'-H₃), 1.26 - 1.35 (10H. m. C3'-H₂, C4'-H₂, C5'-H₂, C6'-H₂, C7'-H₂), 1.51 - 1.65 (2H. m. C2'-H₂), 1.98 - 2.15 (4H. m. C8' - H₂, C17'-H₂), 2.53 (2H. t, J 7.3 Hz. C1'-H₂), 2.66 (2H. t, J 6.7 Hz. C3'-H₂), 2.78 (2H. t, J 7.3 Hz. C1-H₂), 2.66 (2H. t, J 6.7 Hz. C3'-H₂), 2.78 (2H. t, J 6.7 Hz. C2'-H₂), 2.75 - 2.84 (4H. m. C11'-H₂, C14'-H₂) 5.27 = 5.46 (6H. m. C9'-H. C10'-H. C12'-H. C13'-H. C15'-H. C16'-H);
¹³C n.m.r. (50MHz. CDCl₃) δ 178.03s. 131.92d. 130.28d. 128.24d. 127.66d. 127.10d. 34.67t. 32.19t. 29.58t. 29.50t. 29.37t. 29.20t. 29.16t. 28.81t. 27.19t. 26.59t. 25.59t. 25.50t. 20.52t. 14.245q.

Z,Z,Z-(5,8,11,14-Octadecatrienylthio)succinic acid (7a)

[α-carboxymethyl-β-thia 23:4(n-6)]

1 ¹H n.m.r. (300 MHz, CDCl₃) δ 0.89 (3H. t, J 6.8 Hz. C20'-H₃), 1.23 - 1.53 (8H. m. C3'-H₂, C17'-H₂, C18'-H₂, C19'-H₂), 1.56 - 1.70 (2H. m. C2'-H₂), 2.03 - 2.13 (4H. m. C4' - H₂, C16'-H₂), 2.65 - 2.86 (9H. m. C7'-H₂, C10'-H₂, C13'-H₂, C1'-H₂, CHHCO₂H), 3.01 (1H. dd, J 12.1. 17.6 Hz. CHHCO₂H), 3.64 (1H. dd, J 4.0. 12.1 Hz. C2-H), 5.32 - 5.43 (8H. m. C5'-H. C6'-H. C8'-H. C9'-H. C11'-H. C12'-H. C14'-H. C15'-H);

25

SYNTHESIS OF HYDROXY AND HYDOPEROXY DERIVATIVES.

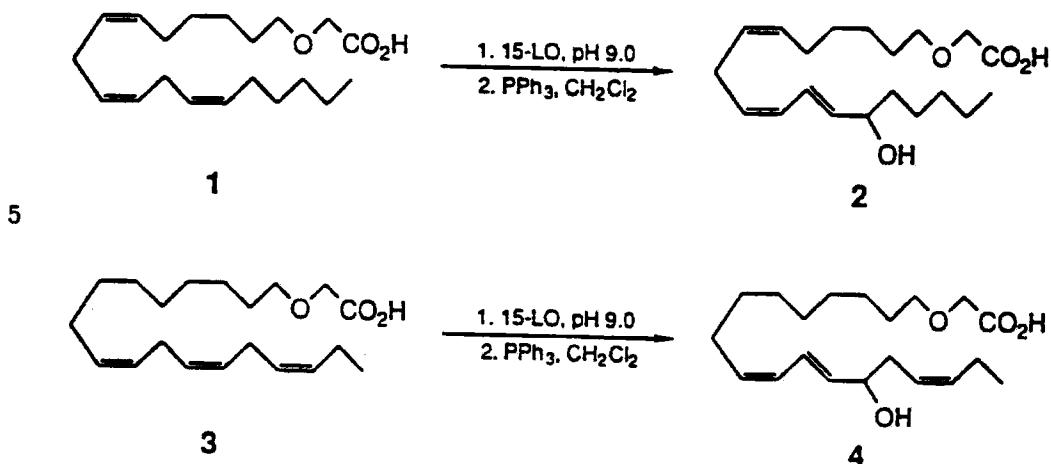
The hydroperoxide derivatives of arachidonic acid (Figs 1b-1g) are obtained separately from enzyme-catalysed reactions of Fig 1a. or as a mixture by autoxidation of Fig 1a. The components of the autoxidation mixture Figs 1b-

30 1g, which vary in ratio depending on the reaction conditions, can be separated by high performance liquid chromatography on silica. Reduction of the hydroperoxides Figs 1b-1g, either separately or as a mixture affords the corresponding alcohols Figs 1h-1m. These can be converted to the corresponding peroxides Fig 1n (R-alkyl or aryl), having the same
 35 substitution pattern as the hydroperoxides Figs 1b-1g and alcohols Figs 1h-1m. by treating with a variety of reagents including the corresponding alkyl

or aryl hypohalites (ROX). Mixtures of either the alcohols Figs 1h-1m or the peroxides Fig 1n can also be separated by high performance liquid chromatography.

- 5 In a similar fashion other naturally occurring unsaturated fatty acids (eg 22:6 (n-3)), and modified fatty acids such as Figs 1o-1x (Y=H) and related compounds prepared by elaboration of acids other than arachidonic acid can be used to prepare hydroperoxy, hydroxy and peroxy derivatives Figs 1o-1x (Y=OOH, OH, OOR), analogous to Figs 1b-1n, where the substitution pattern
- 10 is determined by the allyic oxidation. The acids Fig 1o and Fig 1p (Y=H) can be prepared by aldol condensation of the corresponding aldehyde of Fig 1a and Fig 1p can also be prepared via a Wittig reaction of the same aldehyde or the corresponding halide, while the acid Fig 1q can be prepared via a Wittig reaction of the corresponding C19 aldehyde or halide. The acid Fig 1r can be
- 15 prepared by aldol condensation of a corresponding ester of Fig 1a, while Figs 1s-1x are obtained by ether or thioether synthesis, through nucleophilic substitution or metal catalysed coupling reactions, and the thioethers Figs 1v-1x can be oxidised to the corresponding sulphoxides and sulphones. The amino acid derivatives Fig 1y and Fig 1z can be obtained by coupling the
- 20 corresponding fatty acid (Fig 1a) and glycine and aspartic acid respectively.

Synthesis of Hydroxy β -oxa Fatty Acids



10

REAGENTS

β-oxa Fatty Acids (1,3) - synthesized from the corresponding fatty alcohols by rhodium acetate catalysed coupling with diazoacetate esters.

Triphenylphosphine - Aldrich Chemical Company.

15 *Potassium Dihydrogen Orthophosphate - Ajax Chemicals*

Soybean 15-Lipoxygenase - Aldrich Chemical Company

All solvents were distilled prior to use.

Preparative layer chromatographies were performed on Merck Silica Gel 60 PF₂₅₄ containing gypsum: Art 7749

20

PROCEDURE

Hydroxy β -oxa Fatty Acids 2 and 4

The relevant fatty acid, 1 and 3, (ca. 50 mg) was dissolved in phosphate buffer (0.1 M, pH = 9.0, ca. 45 ml) at 30°C. Soybean 15-lipoxygenase (ca. 8 mg) in phosphate buffer (ca. 5 ml) was added and oxygen was bubbled through the stirred solution for 10 min. Triphenylphosphine (ca. 50 mg) was added in dichloromethane (ca. 50 ml) followed by hydrochloric acid (0.2 M, ca. 20 ml) and the solution was stirred at 0°C for 20 min. The crude reaction

mixture was extracted with dichloromethane and the resulting extract was concentrated under a stream of dry nitrogen. The residue was purified by preparative layer chromatography on silica, eluting with ethyl acetate/hexane/acetic acid (80:20:0.1), to afford the respective hydroxy β -oxa fatty acid, 2 or 4, as an oil

Z, Z, E-(13-Hydroxy-6,9,11-octadecatienyloxy)acetic acid (2)

[16'-OH- β -oxa 21:3(n-6)]

¹H n.m.r. (300 MHz, CDCl₃) δ 0.89 (3H, *t*, *J* 6.8 Hz, C18'-H₃), 1.25 - 5 1.45 (10H, *m*, C3'-H₂, C4'-H₂, C15'-H₂, C16'-H₂, C17'-H₂), 1.59 - 1.73 (2H, *m*, C2'-H₂), 2.01 - 2.12 (4H, *m*, C5' - H₂, C14'-H₂), 2.81 (2H, *t*, *J* 5.8 Hz, C8'-H₂), 3.52 - 3.60 (2H, *m*, C1'-H₂), 4.10 (2H, *s*, C2-H₂), 4.20 (1H, *dt*, *J* 6.0, 6.7 Hz, C13'-H), 5.29 - 5.45 (3H *m*, C6'-H, C7'-H, C9'-H), 5.70 (1H, *dd*, *J* 6.7 15.2 Hz, C12'-H) 5.99 (1H, *dd*, *J* 10.9 Hz, C10'-H) 6.55 (1H, *dd*, *J* 10.9, 15.2 Hz, C11'-H)

10

Z, E, Z-(13-Hydroxy-9, 11, 15-octadecatrienyloxy)acetic acid (4)

[16'-OH- β -oxa 21:3(n-6)]

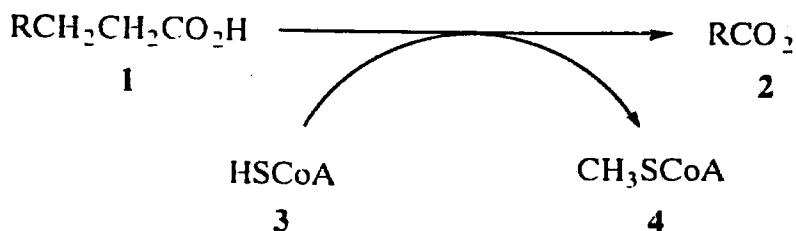
¹H n.m.r. (300 MHz, CDCl₃) δ 0.95 (3H, *t*, *J* 7.6 Hz, C18'-H₃), 1.17 - 15 1.40 (10H, *m*, C3'-H₂, C4'-H₂, C5'-H₂, C6'-H₂, C7'-H₂), 1.48- 1.63 (2H, *m*, C2'-H₂), 1.99 - 2.12 (2H, *m*, C8' - H₂), 2.13 - 2.22 (2H, *m*, C14'-H₂) 2.28 - 2.37 (2H, *m*, C17'-H₂), 3.44 - 3.50 (2H, *m*, C1'-H₂), 3.80 - 3.92 (2H, *s*, C2'-H₂), 4.14 - 4.20 (1H, *dt*, *J* 6.4, 14.9 Hz, C13'-H), 5.30 - 5.60 (3H, *m*, C9'-H, C15'-H, C16'-H), 5.67 (1H, *dd*, *J* 6.4, 14.9 Hz, C12'-H), 5.95 (1H, *dd*, *J* 11.1, 11.1 Hz, C10'-H), 6.49 (1H, *dd*, *J* 11.1, 14.9 Hz C11'-H)

20

Effect of novel fatty acids on the action of Acyl-CoA-Oxidase.

β -oxidation of fatty acids

5 β -oxidation is the main oxidative metabolic fate of fatty acids (1), the net process being characterized by the degradation of the fatty acid carbon chain by two carbon atoms with the concomitant production of equimolar amounts of acetyl-coenzyme A (4) (Scheme 1).



Scheme 1

10 Acyl-CoA-oxidase

The first step in the β -oxidation pathway is controlled by acyl-CoA oxidase. In peroxisomes, acyl-CoA oxidase catalyses the reaction of acyl-coenzyme A thioesters with oxygen to produce enoyl-coenzyme A thioesters. In the presence of peroxidase enzyme the concomitant production of hydrogen peroxide is cycled back to oxygen with the net generation of one mole of water for each mole of acyl-coenzyme A thioester oxidized.

Novel fatty acids

20 Fatty acids bearing a blocking group at the β -position were anticipated as being inert to the usual course of β -oxidation. Novel fatty acids thus bearing a heteroatom at the β -position have been synthesised. The compounds used in the current investigation were arachidonyloxyacetic acid [β -oxa 23:4(n-6)], gamma linolenyloxyacetic acid [β -oxa 21:3(n-6)], linolenyloxyacetic acid [β -oxa 21:3(n-3)] and arachidonylthioacetic acid [β -thia 23:4(n-6)]. The oxidation of arachidonyl-thiopropionic acid [γ -thia 22:4(n-6)] was also 25 investigated in order to assess the effect of a blocking group at the γ -position of a fatty acid upon β -oxidation.

Acyl-CoA Oxidase assay

Acyl-CoA oxidase from *Arthrobacter* species is available from Boehringer Mannheim in stabilized tablet form and is commonly used together with acyl-CoA synthetase for the measurement of free acids in serum or plasma.

5 Assay principle

Free fatty acids are, in the presence of the enzyme acyl-CoA synthetase (acyl CS), converted by adenosine-5'-triphosphate (ATP) and coenzyme A (CoA) into acyl-coenzyme A (acyl-CoA.) resulting in adenosine-5'-monophosphate (AMP) and pyrophosphate being produced.

10 Acyl-CoA reacts with oxygen (O₂) in the presence of acyl-CoA-oxidase (ACOD) to form 2.3-enoyl-coenzyme A (enoyl-CoA.) with the generation of hydrogen peroxide.

The resulting hydrogen peroxide (H₂O₂) converts 4-aminoantipyrine (4-AA) and 2.4.6-tribromo-3-hydroxybenzoic acid (TBHB) into a red dye in the

15 presence of peroxidase (POD). The dye is measured in the visible region at 546 nm.

Assay procedure

Wavelength: 546 nm

Micro Cuvette: 1 cm light path

20 Temperature: 25°C (± 1°C)

Assay volume: 1.15 ml

Measurement of reaction mixtures against air

Pippetted into cuvette:

1.00 ml reaction mixture: KH₂PO₄ (87 mmol/l), Mg²⁺ (1.3 mmol/l),

25 ATP (1.5 mmol/l), CoA (0.26 mmol/l), POD (21.7 kU/l), Acyl CS (165 U/l).

0.05 ml sample solution of free fatty acid(s).

[for blank assay 0.05 ml of distilled water was substituted]

Sample was mixed and temperated at 25°C for approx. 10 min.

30 Following this, addition of:

0.05 ml N-ethyl maleinamide solution: (0.86 mmol/l)

Absorbance (A1) of solution was measured at 546 nm followed by the addition of:

0.05 ml ACOD solution: (8.7 kU/l).

At t = 10 seconds after the addition of ACOD to the reaction mixture the 5 change in visible absorption of the solution at 546 nm was measured for 20 min.

Assay calibration / authentication

Sample concentration dependance

Arachidonic acid (20:4(n-6). 10) was assayed at the following concentrations:

10 4.3 mM. 2.1 mM. 0.99 mM. 0.49 mM. 0.25 mM and 0 mM (blank assay).

As per *Figure 1* the assay shows rate dependance on the concentration of fatty acid.

Enzyme (ACOD) concentration dependance

A sample solution of arachidonic acid (20:4(n-6). 10) at a concentration of 15 0.99 mM was assayed using acyl-CoA oxidase concentrations as follows: 200 kU/l. 100 kU/l and 50 kU/l.

As per *Figure 2* the assay shows rate dependance on the concentration of enzyme (ACOD).

Assay of novel fatty acids

20 Solutions of the novel compounds at concentrations of 1.0 mM, 1.0 mM, 1.0 mM, 1.0 mM and 1.1 mM respectively were assayed using an acyl-CoA oxidase concentration of 100 kU/l.

As per *Figures 3 – 6* the assay indicates that the β -oxa compounds and the β -thia compounds respectively, are not substrates for acyl-CoA oxidase (Figure 25 11A).

Effect of novel compounds on arachidonic acid metabolism by Acyl-CoA oxidase

Sample solutions containing equimolar concentrations of arachidonic acid (10) and each of the novel compounds were assayed using an acyl-CoA oxidase concentration of 100 kU/l.

The assay shows that the novel compounds have no effect on the uptake of arachidonic acid by acyl-CoA oxidase.

5 Further samples containing novel compounds and arachidonic acid (10) at relative concentrations of 4:1 and 16:1 were assayed and indicated no appreciable effect on the uptake of arachidonic acid by acyl-CoA oxidase (not shown).

10 Covalent Coupling of Fatty Acids to Amino Acids

Polyunsaturated fatty acid (1.80 mmol), HOSu (0.41 g, 3.60 mmol) and the amino acid t-butyl ester (3.60 mmol) were dissolved together in dimethylformamide (3 mL) and the mixture cooled in ice bath.

15 Dicyclohexylcarbodiimide (0.44 g, 2.16 mmol) in DMF (0.3 mL) and N-methylmorpholine (0.73 g, 7.20 mmol) were added. The mixture stirred for 20 hours, after which time some unreacted polyunsaturated fatty acid remained. More DCC (0.10 g) was added and the mixture stirred for further 20 hours. DCU was filtered off and the product isolated by reversed phase

20 HPLC. The purified product was concentrated to an oil and trifluoroacetic acid (30 mL) as added. After an hour stirring, the trifluoroacetic acid was evaporated to leave the product as a brown oil which was redissolved in CH₃CN (6 mL) and purified by reversed phase HPLC. The pure fractions obtained were combined, concentrated and lyophilised (in t-butanol) to

25 leave the product as a brown oil.

TABLE 1
DIRECT ANTIMALARIAL ACTIVITY OF NOVEL FATTY ACIDS

COMPOUND	EC50 (μ M)
β -oxa 21:3 (n-6)	<30
β -oxa 21:3 (n-3)	30
β -oxa 21:4 (n-3)	40
β -thia 21:3 (n-6)	30-50
β -thia 21:3 (n-3)	30-35
β -oxa 25:6 (n-3)	<60
γ -thia 22:0 (n-6)	45
γ -thia 22:3 (n-6)	45
β -oxa 23:4 (n-6)	<30
β -thia 23:4 (n-6)	15-30
γ -thia 24:4 (n-6)	15
β -thia 25:6 (n-3)	25
α - carboxymethyl- β -thia 23:4 (n-6)	50
16-OH- β -oxa 21:3 (n-6)	50
16-OH- β -oxa 21:3 (n-3)	40

TABLE 2
FATTY ACID STIMULATED NEUTROPHIL-MEDIATED ANTI-MALARIAL
ACTIVITY

COMPOUND	% INHIBITION RELATIVE TO 22:6 AT 16 μ M
β -oxa 21:3 (n-3)	45
β -oxa 21:4 (n-3)	85
β -thia 21:3 (n-6)	85
β -thia 21:3 (n-3)	73
β -thia 23:4 (n-6)	150
γ -thia 24:4 (n-6)	91
β -thia 25:6 (n-3)	47
16-OH- β -oxa 21:3 (n-6)	33

TABLE 3
EFFECT OF MP8, MP11, MP14 ON K1 CHLOROQUINE RESISTANT
P.FALCIPARUM

COMPOUND	% GROWTH INHIBITION
CHLOROQUINE	12.4
β -thia 23:4 (n-6)	97
γ -thia 24:4 (n-6)	97
β -thia 25:6 (n-3)	98

TABLE 4
EFFECT OF FATTY ACIDS ON MITOGEN-INDUCED PERIPHERAL BLOOD
MONONUCLEAR CELL PROLIFERATION

COMPOUND	% INHIBITION OF INDUCED PROLIFERATION AT 20 μ M PUFA
β -oxa 21:3 (n-6)	17
β -thia 21:3 (n-3)	28
β -thia 25:6 (n-3)	97
16-OH- β -oxa 21:3 (n-6)	50

TABLE 5
EFFECT OF PUFA ON PHA-INDUCED TNF α PRODUCTION

COMPOUND	% INHIBITION OF CYTOKINE PRODUCTION (PUFA AT 20 μ M)
β -oxa 21:3 (n-6)	38
β -oxa 21:3 (n-3)	39
β -thia 21:3 (n-6)	17
β -thia 21:3 (n-3)	17
γ -thia 22:3 (n-6)	41
γ -thia 22:3 (n-3)	25
β -oxa 23:4 (n-6)	25
β -thia 23:4 (n-6)	35
γ -thia 24:4 (n-6)	34
β -thia 25:6 (n-3)	90
16-OH- β -oxa 21:3 (n-6)	71
16-OH- β -oxa 21:3 (n-3)	68

TABLE 6
EFFECT OF PUFA ON STAPH AUREUS INDUCED INTERFERON γ PRODUCTION BY PERIPHERAL BLOOD MONONUCLEAR CELLS

COMPOUND	% INHIBITION (PUFA AT 20 μ M)
β -oxa 21:3 (n-6)	89
β -thia 21:3 (n-6)	45
β -oxa 25:6 (n-3)	44
β -oxa 23:4 (n-6)	89
β -thia 23:4 (n-6)	64
β -thia 25:6 (n-6)	96
16-OH- β -oxa 21:3 (n-6)	77
16-OH- β -oxa 21:3 (n-3)	65

CLAIMS:-

1. A polyunsaturated fatty acid compound having antimalarial and/or neutrophil stimulatory activity, the polyunsaturated fatty acid containing 18-25 carbons and 1-6 double bonds and wherein the polyunsaturated fatty acid compound has one or two substitutions selected from the group consisting of β oxa, γ oxa, β thia and γ thia.
2. A polyunsaturated fatty acid compound as claimed in claim 1 in which the polyunsaturated fatty acid compound includes a further substitution selected from the group consisting of hydroxy, hydroperoxy, peroxy, carboxymethyl substitutions or attached to an amino acid.
3. A polyunsaturated fatty acid compound as claimed in claim 1 or claim 2 in which the polyunsaturated fatty acid compound contains 20-25 carbon atoms and 3-6 double bonds.
4. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 3 in which the polyunsaturated fatty acid compound is an n-3 to n-6 fatty acid.
5. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 21 carbons with 3-4 double bonds containing a β oxa or β thia substitution.
6. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 22 carbon atoms with 3-4 double bonds containing a γ thia or β oxa substitution.
7. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 23 carbons with 3-4 double bonds containing a β thia substitution.
8. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 24 carbons with 3-4 double bonds containing a γ thia substitution.
9. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 25 carbons with 3-6 double bonds containing a β oxa substitution.
10. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is 25 carbons with 3-6 double bonds containing a β thia substitution.

11. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 10 in which the polyunsaturated fatty acid compound has a ω hydroxy substitution.
12. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound has 23 carbons, 3-6 double bonds, β thia and α -carboxymethyl group.
13. A polyunsaturated fatty acid compound as claimed in any one of claims 1 to 4 in which the polyunsaturated fatty acid compound is covalently attached via the carboxy terminus to an amino acid.
14. A polyunsaturated fatty acid compound as claimed in claim 13 in which the polyunsaturated fatty acid compound is covalently attached to aspartic acid or glycine.
15. A method of producing an unsaturated oxa substituted fatty acid comprising reacting an unsaturated fatty acid alcohol with a carbene such that the carbene is inserted in the O-H bond of the alcohol.
16. A method as claimed in claim 15 in which the unsaturated fatty acid alcohol contains 18-25 carbon atoms and 1-6 double bonds.
17. A method as claimed in claim 15 or claim 16 in which the carbene is synthesized via rhodium acetate catalysed of a diazo compound.
18. A method as claimed in any one of claims 15 to 17 in which an unsaturated β oxa substituted fatty acid is produced.
19. A method of treating inflammation in a subject. the method comprising administering to the subject a therapeutically effective amount of an anti-inflammatory composition comprising at least one hydroxy, hydroperoxy, or peroxy derivative of a polyunsaturated fatty acid having a carbon chain of 18 to 24 carbon atoms and having 1-6 cis or trans double bonds.
20. A method as claimed in claim 18 in which the poly unsaturated fatty acid contains oxygen or sulphur atoms within the carbon chains as oxa or thia derivatives.
21. A method as claimed in claim 18 in which the polyunsaturated fatty acid is selected from the group consisting of the C20:4n-6 (5,8,11,14-eicosatetraenoic acid), C20:5n-3 (5,8,11,14,17-eicosapentaenoic), C22:6n-3 (4,7,10,13,16,19-docosahexaenoic acid) and arachidonic acid.

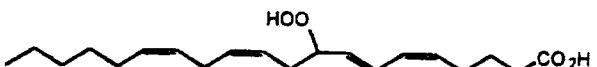
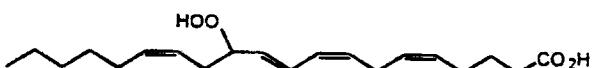
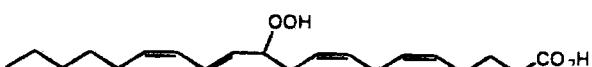
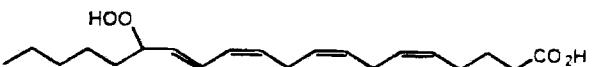
**Fig 1a****Fig 1b****Fig 1c****Fig 1d****Fig 1e****Fig 1f****Fig 1g****Fig 1**



Fig 1h

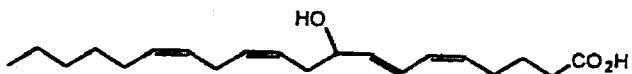


Fig 1i

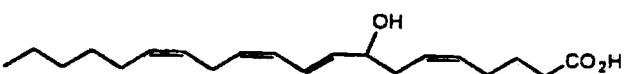


Fig 1j



Fig 1k

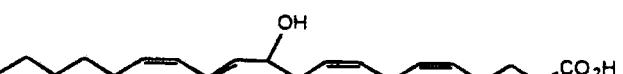


Fig 1l

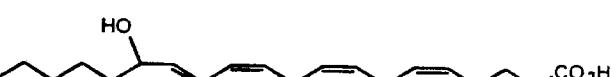


Fig 1m

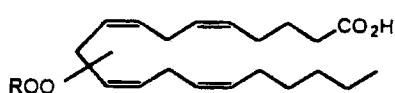


Fig 1n

Fig 1 (cont)

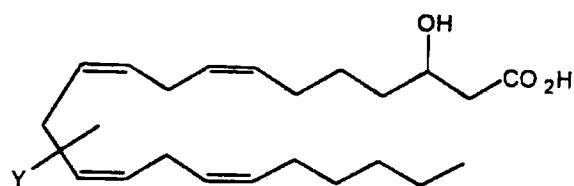


Fig 1o

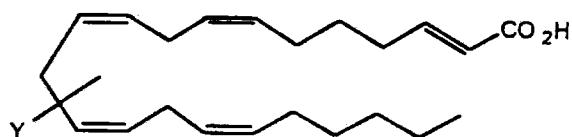


Fig 1p

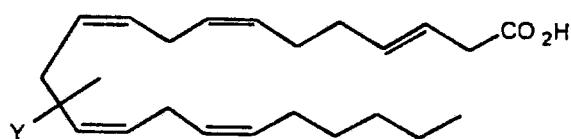


Fig 1q

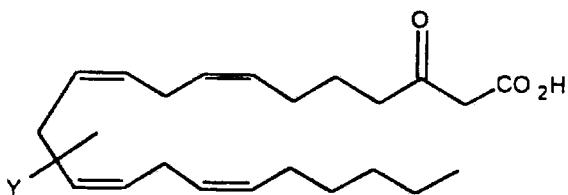


Fig 1r

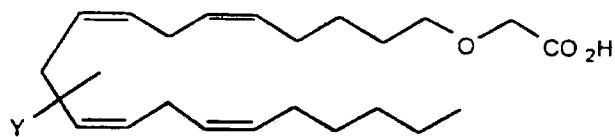


Fig 1s

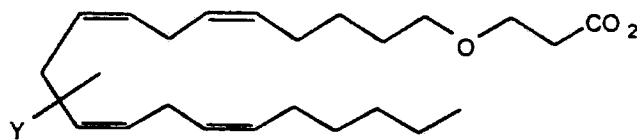
Fig 1t
SUBSTITUTE SHEET (RULE 26)

Fig 1 (cont)

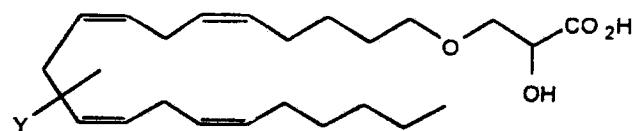


Fig 1u

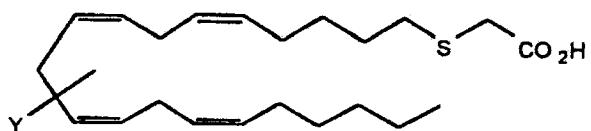


Fig 1v

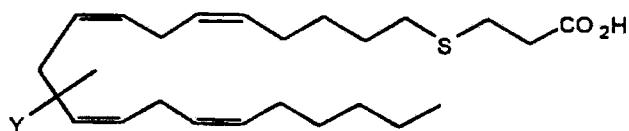


Fig 1w

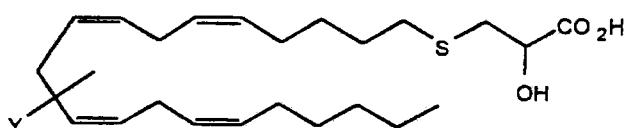


Fig 1x

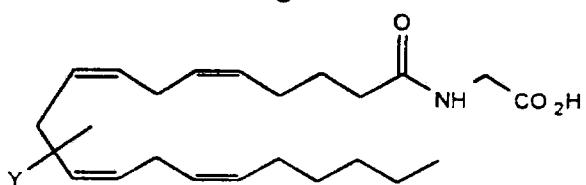


Fig 1y

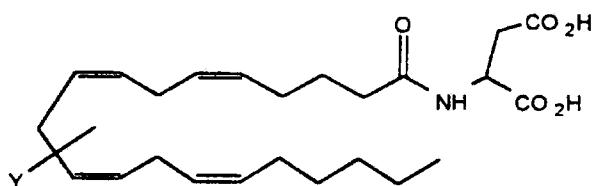


Fig 1z

FIG 1 (cont)

FIG. 2



Fig. 2a



Fig. 2b



Fig. 2c



Fig. 2d



Fig. 2e

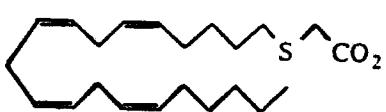


Fig. 2f

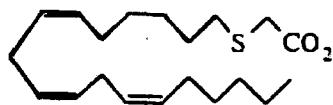


Fig. 2g

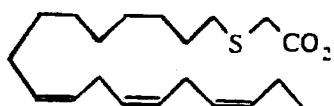


Fig. 2h



Fig. 2i



Fig. 2j

FIG. 2 (cont'd.)



Fig. 2k

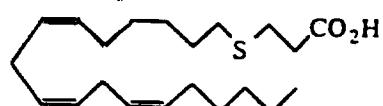


Fig. 2l

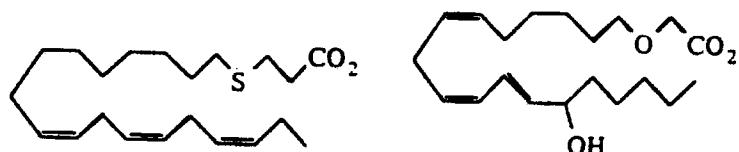


Fig. 2m

Fig. 2n

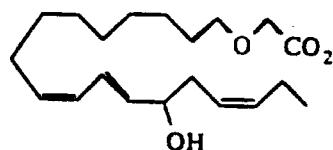


Fig. 2o

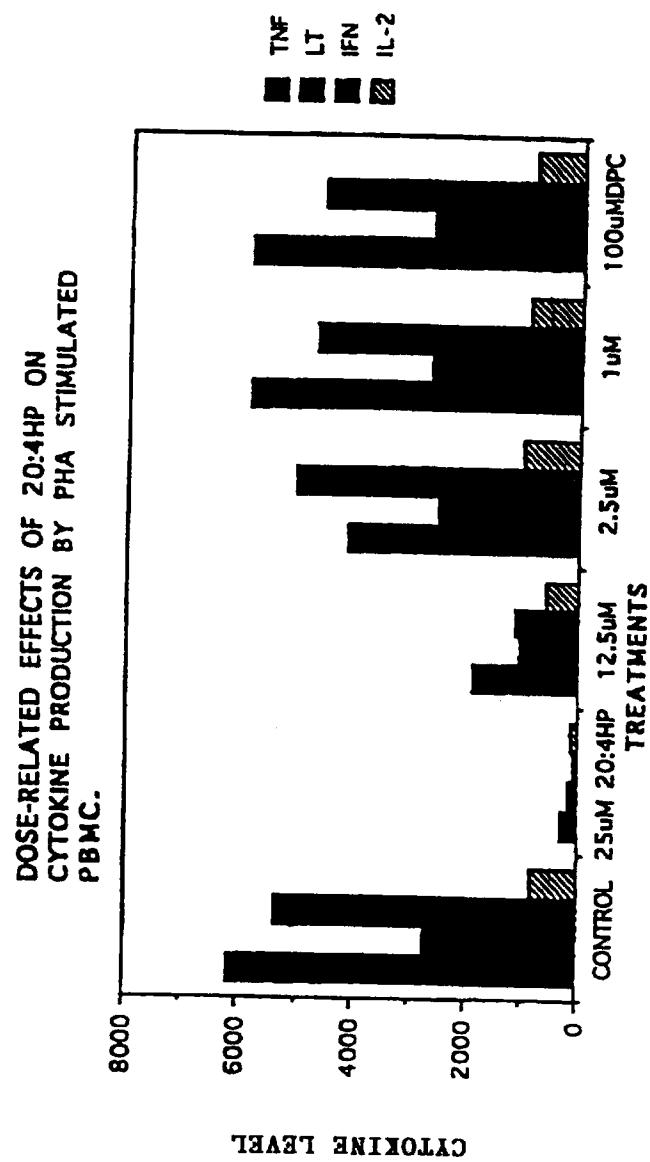


Fig. 3

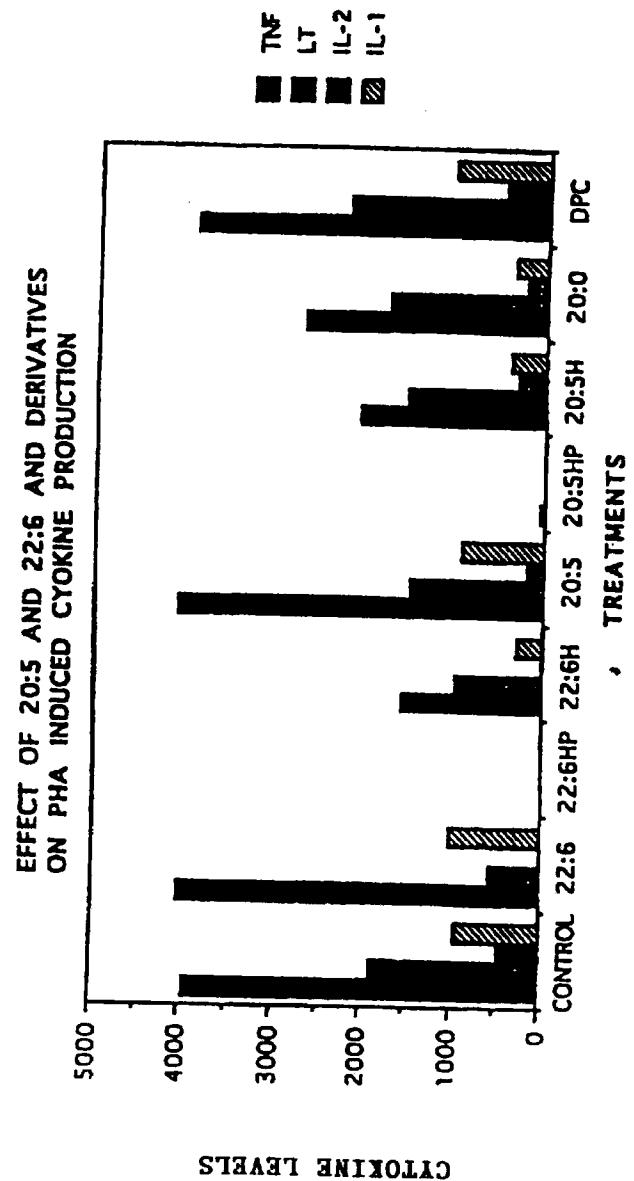


Fig. 4

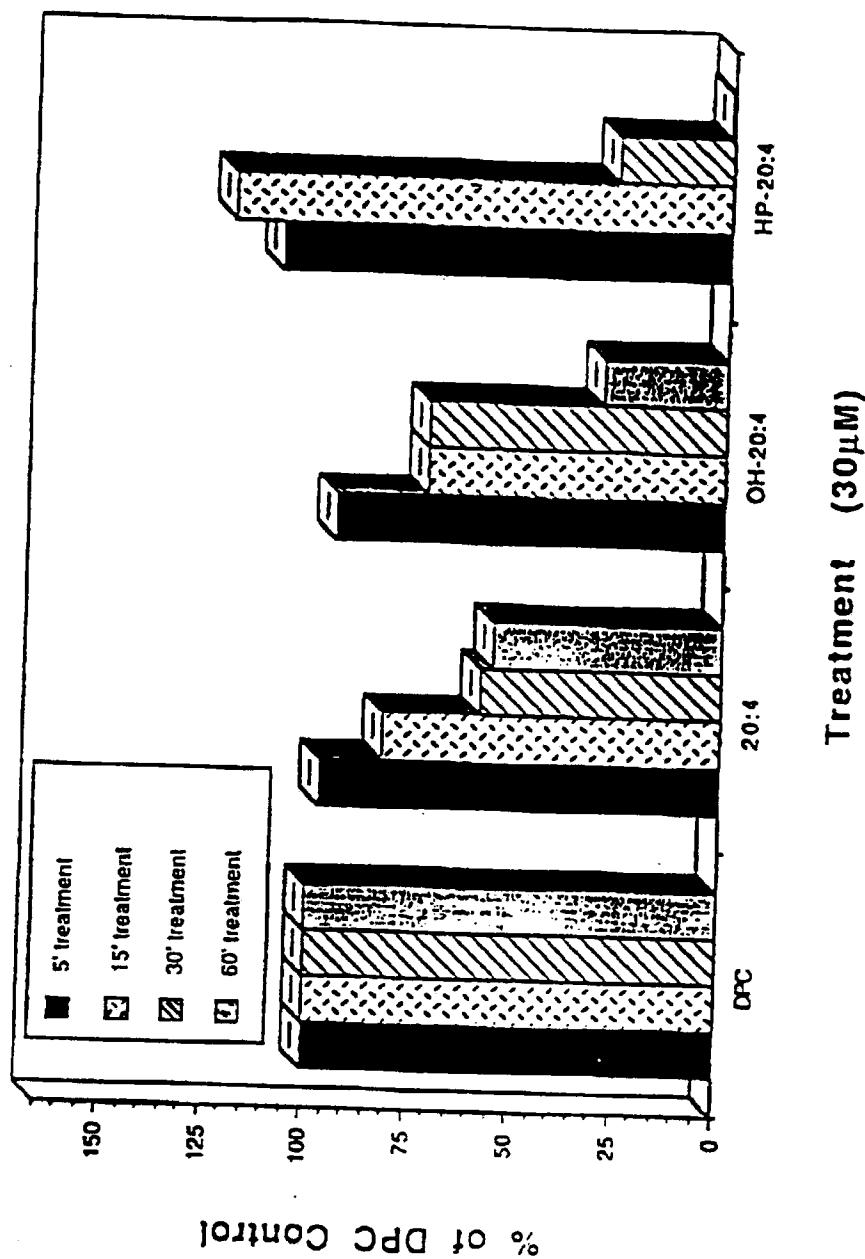
Effect of FAs on the Expression of E-selectin on HUVEC

FIG. 5

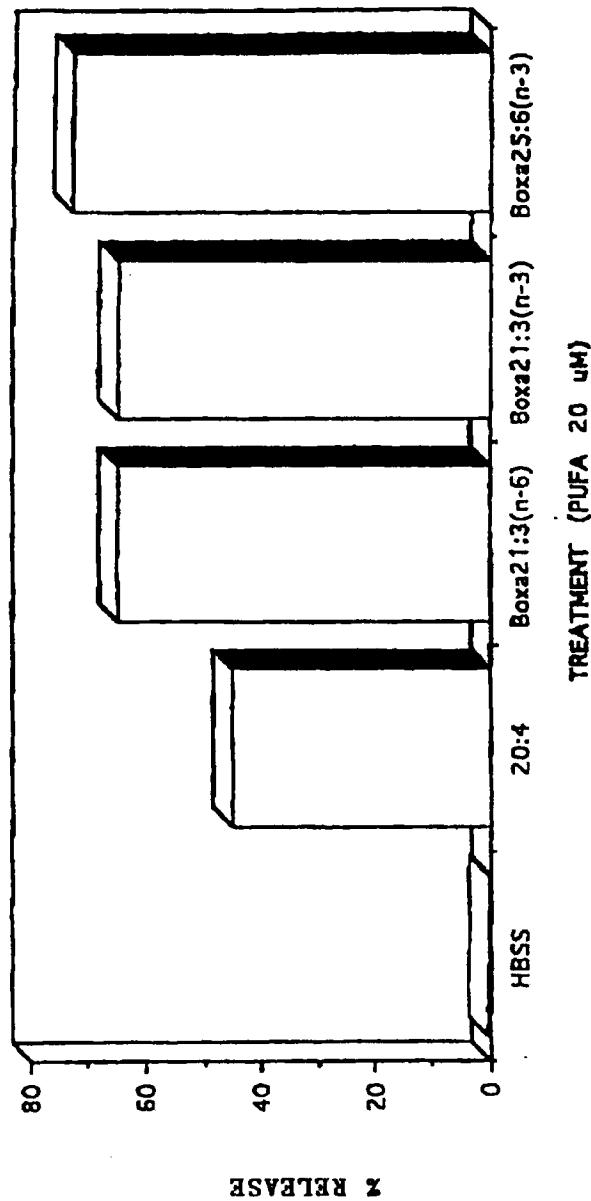


FIG. 6.

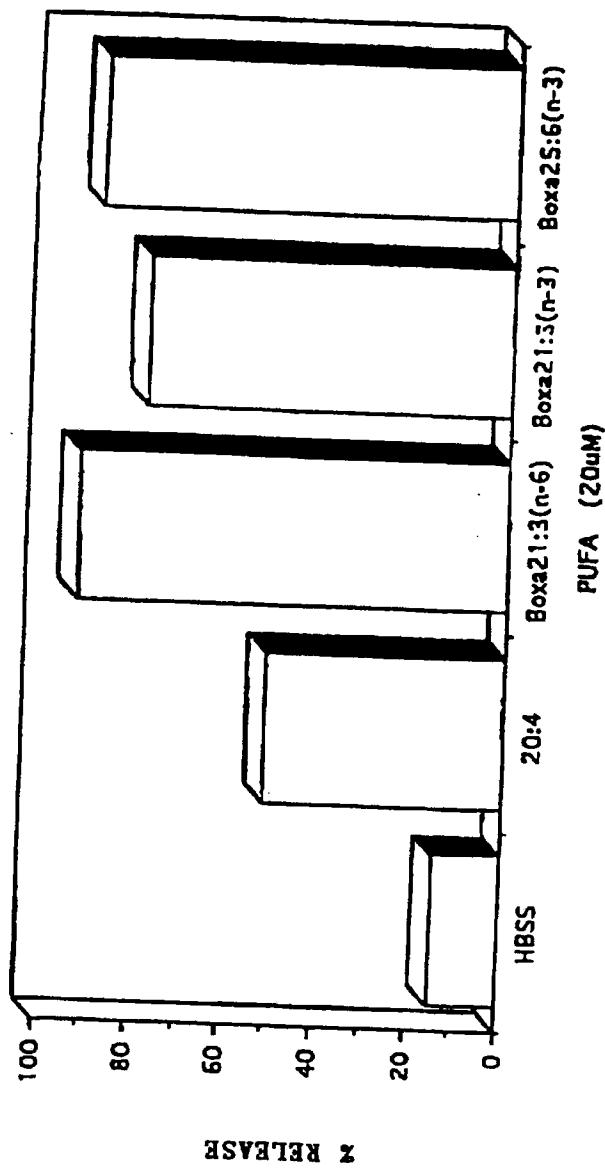


Fig. 7

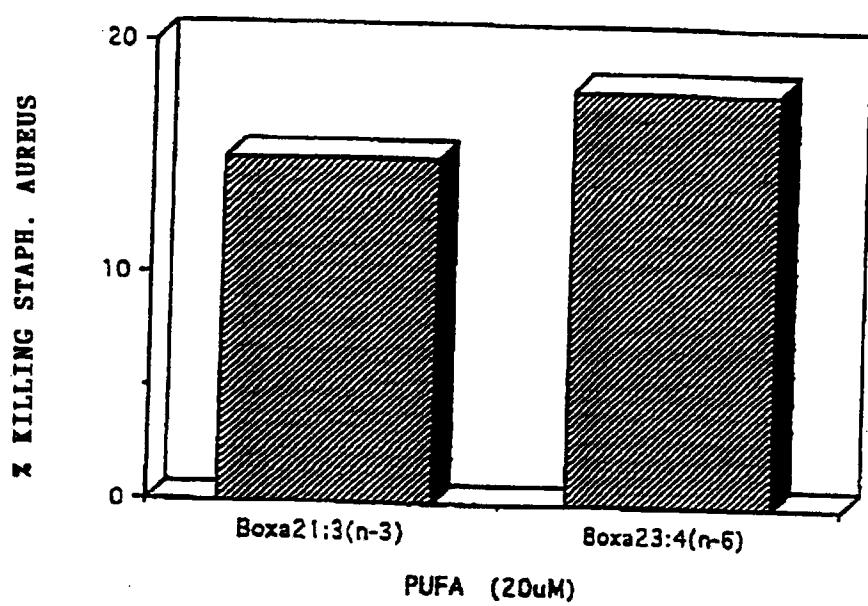


FIG. 8

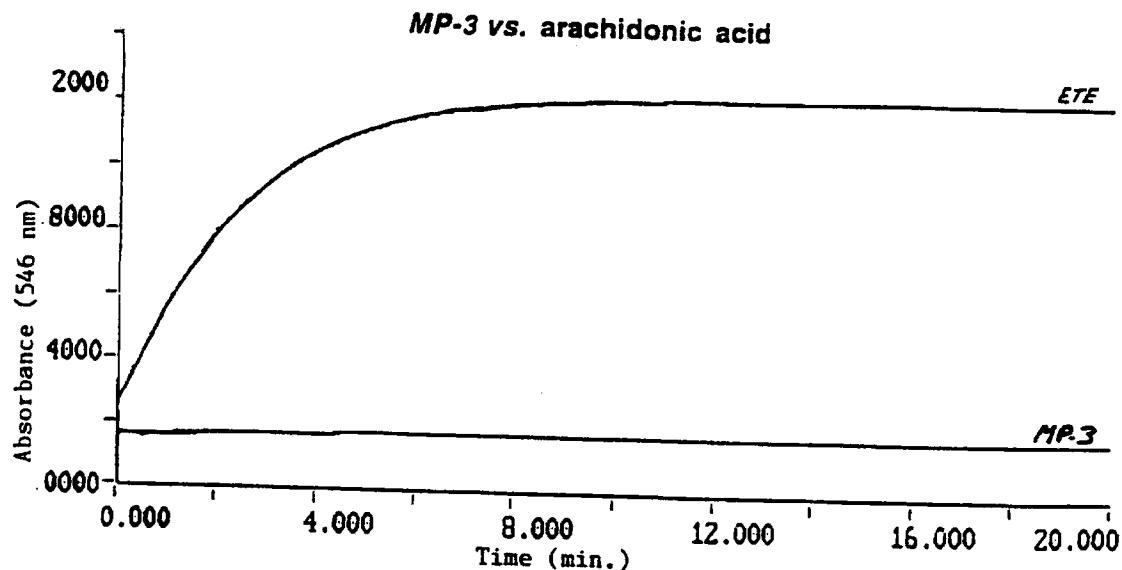


FIG. 9a

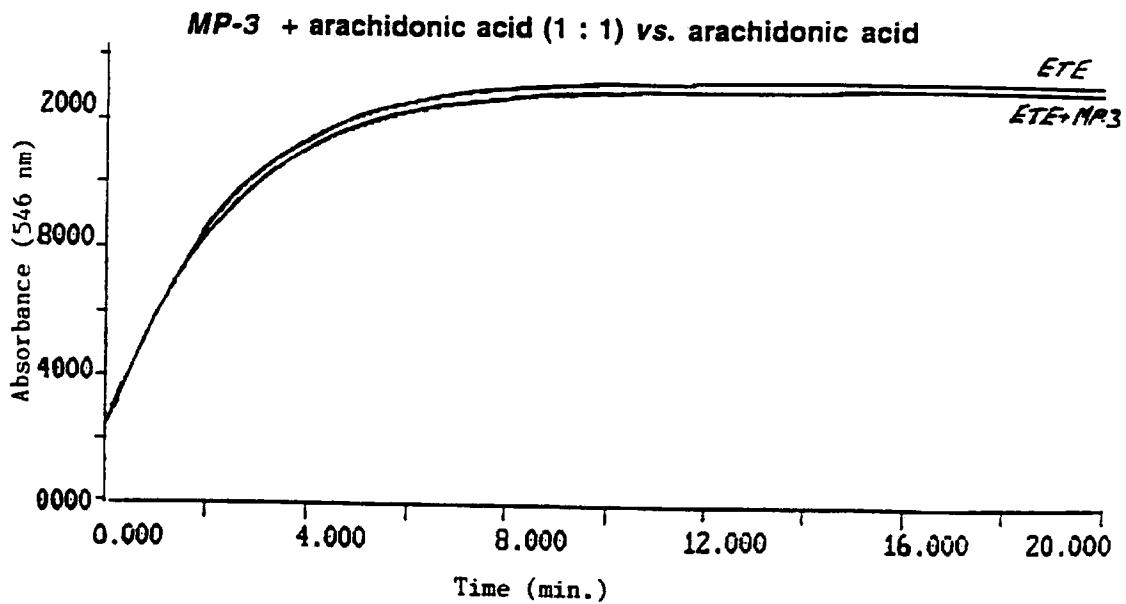


FIG. 9b

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/AU 95/00677

A. CLASSIFICATION OF SUBJECT MATTER

Int Cl⁶ C07C 323/52 59/60 409/42 409/16 409/04 51/367 A61K 31/20 31/19 31/16 // C07C 69/734

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07C 59/60 149/20 179/06 179/02 179/22 179/025 59/27 323/52 409/04 409/16 409/42 51/367 51/36
A61K 31/20 31/19

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
AU: IPC as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

DERWENT: fatty()acid: or eicosa: or hydroperoxy: or peroxy: or carbene: or hydroxy:

CHEMICAL ABSTRACTS: fatty()acid: or eicosa:

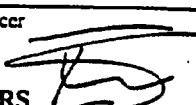
C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
X	EP 345038 A2 (NORSK HYDRO A.S.) 6 December 1989 entire document	1, 3-7, 9, 10
X	EP 260655 A2 (SUMITOMO PHARMACEUTICALS CO. LTD.) 23 March 1988 entire document	18, 20
X	Journal of Lipid Research, volume 31, No. 5, 1990, C. von Schacky et al. "Platelet-neutrophil interactions 12S, 20- and 5S., 12S-dihydroxyeicosapentaenoic acids: two novel neutrophil metabolites from platelet-derived 12S-hydroxyeicosapentaenoic acid", pages 801-810. entire document	18, 20

Further documents are listed in the continuation of Box C

See patent family annex

• Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier document but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	"&" document member of the same patent family

Date of the actual completion of the international search 8 January 1996	Date of mailing of the international search report 18.01.96
Name and mailing address of the ISA/AU AUSTRALIAN INDUSTRIAL PROPERTY ORGANISATION PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No.: (06) 285 3929	Authorized officer  T. SUMMERS Telephone No.: (06) 283 2291

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/AU 95/00677

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Chemical Abstracts, volume 119, No. 19, issued 8 November 1993, R. Bunsho et al, "Anti-inflammatory agents containing [5Z, 8Z, 11Z, 13E, 15(S), 17Z]-15-hydroxyeicosapentaenoic acid derivatives and having immune-modulating effect", page 87, column 2, abstract no. 195671a. abstract	18, 20
A	WO 95/09622 A (PEPTIDE TECHNOLOGY LTD et al), 13 April 1995 entire document	1-20
A	WO 93/00084 A (ADELAIDE CHILDREN'S HOSPITAL), 7 January 1993 entire document	1-20
A	Interleukins, Lymphokines and Cytokines, proceedings from the third International Lymphokine Workshop, 1982, Yael Bromberg et al. "Oxidative metabolism of lymphokine activated macrophages-free fatty acids as second messenger of superoxide generation", pages 561-567. entire document	1-20
P, X	Journal of Clinical Investigation, volume 89, no. 3, 1992, L. M. Kumarulake et al, "Antimalarial properties of n-3 and n-6 polyunsaturated fatty acids: in vitro effects on Plasmodium falciparum and in vivo effects on P. berghei", pages 961-967. entire document	18
A	WO 94/22848 A (PACE-ASCIAK), 13 October 1994 entire document	1-20
A	Biochimica et Biophysica Acta, 1199, 1994, A. Demoz et al. "Modulation of plasma and hepatic oxidative status and changes in plasma lipid profile by n-3 (EPA and DHA), n-6 (corn oil) and a 3-thia fatty acid in rats", pages 238-244. entire document	1-20
A	Proceedings Scandinavian Symposium Lipids, 16th, 1991, S. Skrede et al, "3-Thia fatty acids, clofibrate and omega-3 fatty acids - compounds reducing serum lipids. They might act hypolipidaemic by similar mechanisms", pages 254-270. entire document	1-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No.
PCT/AU 95/00677

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
EP	345038	CA	1329550	DK	2676/89	US	5093365
EP	260655	JP	63072648				
WO	9509622	AU	78482/94				
WO	9300084	AU	21726/92	EP	591303		
WO	9422848	AU	65117/94				

END OF ANNEX